

This Page Is Inserted by IFW Operations  
and is not a part of the Official Record

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning documents *will not* correct images,  
please do not report the images to the  
Image Problem Mailbox.**

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
8 February 2001 (08.02.2001)

PCT

(10) International Publication Number  
**WO 01/08727 A1**

(51) International Patent Classification<sup>7</sup>: **A61M 5/145**

(21) International Application Number: **PCT/US00/20623**

(22) International Filing Date: **28 July 2000 (28.07.2000)**

(25) Filing Language: **English**

(26) Publication Language: **English**

(30) Priority Data:  
**09/365,285** **30 July 1999 (30.07.1999)** **US**

(71) Applicant: **MEDRAD, INC.** [US/US]; One Medrad Drive, Indianola, PA 15051 (US).

(72) Inventors: **NOLAN, William, J., Jr.**; 37 Victory Way, Indianola, PA 15051 (US). **SPOHN, Michael, A.**; 112 Pine Ridge Lane, Butler, PA 16002 (US). **HAURY, John, A.**; 2514 Nicholson Road, Sewickly, PA 15143 (US). **MC-COPPIN, Anthony, S.**; 830 Pittsburgh Road, Butler, PA 16002 (US). **LAZZARO, Frank, A.**; 135 Lucille Drive,

Pittsburgh, PA 15235 (US). **ASHCRAFT, Robert, J., Jr.**; 704 Three Degree Road, Butler, PA 16002 (US). **DEDIG, James, Albert**; 7 Elmhurst Road, Pittsburgh, PA 15220 (US). **BLUE, Christopher, J.**; 2127 Ridge Road, Boston, PA 15135 (US). **BERRY, David, H.**; R.D. 2, Box 57B, Worthington, PA 16262 (US).

(74) Agent: **BRADLEY, Gregory, L.**; Medrad, Inc., One Medrad Drive, Indianola, PA 15051 (US).

(81) Designated States (*national*): **CN, JP.**

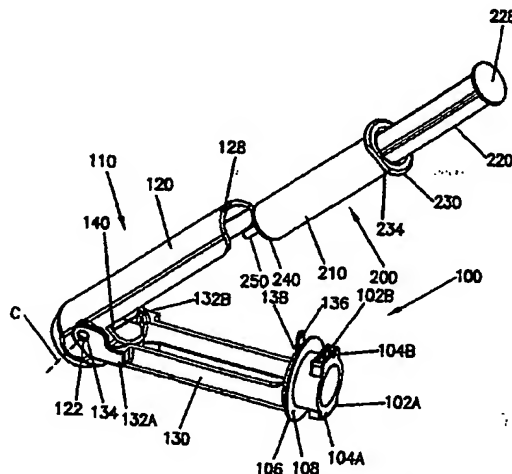
(84) Designated States (*regional*): European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).

Published:

— *With international search report.*

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: **INJECTOR SYSTEMS AND SYRINGE ADAPTERS FOR USE THEREWITH**



(57) Abstract: An adapter includes a syringe carrier adapted to seat at least a portion of the syringe. The syringe carrier includes at least one rearward facing abutment member to abut at least one forward facing surface of a syringe. The syringe carrier includes an opening therein to allow a drive member of an injector to communicate forward force to the plunger through abutment without connective engagement between the drive member and the plunger. The adapter further includes a releasable mounting mechanism positioned to the rear of the syringe carrier to mount the adapter in a desired position relative to the front wall of the injector. An adapter includes a first section and a second section that are rotatable relative to each other about a hinge axis generally perpendicular to a longitudinal axis of the adapter. An adapter includes a first section and a second section that are generally the same in construction, the first section and the section being connectable to form a syringe carrier to seat at least a portion of the syringe.

WO 01/08727 A1

-1-

## INJECTOR SYSTEMS AND SYRINGE ADAPTERS FOR USE THEREWITH

### Field of the Invention

5           The present invention relates to powered injector systems and syringe adapters for use therewith.

### Background of the Invention

10           A number of injector-actuated syringes and powered injectors for use in medical procedures such as angiography, computed tomography, ultrasound and NMR/MRI have been developed. U.S. Patent No. 4,006,736, for example, discloses an injector and syringe for injecting fluid into the vascular system of a human being or an animal. Typically, such injectors comprise drive members such as pistons that connect to a syringe plunger. For example, U.S. Patent No. 4,677,980, the disclosure of which is incorporated herein by reference, discloses an angiographic injector and syringe wherein  
15           the drive member of the injector can be connected to, or disconnected from, the syringe plunger at any point along the travel path of the plunger via a releasable mechanism. A front-loading syringe and injector system is also disclosed in U.S. Patent No. 5,383,858, the disclosure of which is incorporated herein by reference.

20           As discussed in U.S. Patent No. 5,383,858, a syringe used with a front-loading injector preferably includes a readily releasable mounting mechanism for securing the syringe to the front wall of the injector. The use of specifically designed mounting mechanisms, however, prevents the use of syringes of other various types with front-loading injectors. Such syringes may, for example, include a syringe body, a plunger reciprocally mounted therein, and a plunger extension for transfer of force  
25           to the plunger.

U.S. Patent No. 5,520,653, the disclosure of which is incorporated herein by reference, discloses several adapters designed to allow the use of various syringes with a front-loading injector. In one embodiment, the adapter of U.S. Patent No. 5,520,653 includes a syringe carrier having a front end, a rear end, and syringe retaining channel located between the carrier front and rear ends for engaging at least a portion of the syringe flange. Mounting flanges near the rearward end of the carrier releasably mount the carrier in a desired position relative to the front wall of the injector. The adapter of U.S. Patent No. 5,520,653 further includes a follower reciprocally mounted within the carrier. The follower has a front end that engages the syringe plunger extension when the syringe is installed in the carrier. A drive head opening in the carrier communicates with a pair of drive head slots positioned near the rear end of the follower for releasably mounting the follower in a desired position relative to the drive head of the injector.

Although U.S. Patent No. 5,520,653 is a substantial improvement in the art, it remains desirable to develop improved adapters for use with syringes of various types to permit use of such syringes with front-loading injectors.

### **Summary of the Invention**

In general, the present invention provides an adapter for releasably mounting a syringe in a desired position relative to a front-loading powered injector. The syringe includes a body and a plunger slideably positioned within the body. The injector includes a front wall, an opening formed in the front wall, and a drive member reciprocally mounted in the injector. The adapter preferably includes generally a syringe carrier adapted to seat at least a portion of the syringe. The syringe carrier includes at least one rearward facing abutment member to abut at least one forward facing surface on the syringe. The syringe carrier includes an opening therein to allow the drive member of the injector to communicate forward force to the plunger through abutment without connective engagement between the drive member and the plunger. The adapter further includes a releasable mounting mechanism

positioned to the rear of the syringe carrier to mount the adapter in a desired position relative to the front wall of the injector.

The syringe may further include a transition region over which the radius or width of the syringe decreases (for example, a generally frusto-conical region) attached to a forward end of the body. The abutment member may abut a forward facing surface created by the transition region. Preferably, the abutment member abuts the transition region only in the vicinity of the transition from the body to the generally frusto-conical region (for example, at to the outer edge of the transition region).

The syringe may further include a syringe flange attached to a rearward end of the body of the syringe. The abutment member may abut a forward facing surface of the syringe flange. Preferably, the abutment member abuts the syringe flange only in the vicinity of the transition from the body to the syringe flange.

In one embodiment, the adapter includes a first section and a second section rotatable relative to each other about a hinge axis generally perpendicular to a longitudinal axis of the adapter. The first section and the second section are preferably rotatable about the hinge axis to an open position to allow loading of the syringe into the adapter from a position to the rear of the hinge axis. The first section and the second section are also preferably rotatable about the hinge axis to a closed position to form the syringe carrier.

In another embodiment, the adapter includes a first section and a second section that are generally the same in construction. The first section and the second section are connectable to form the syringe carrier and the releasable mounting mechanism.

The present invention also provides an adapter for releasably mounting a syringe in a desired position relative to a powered injector. The syringe preferably

includes a body and a plunger slideably positioned within the body as discussed above. The injector preferably includes a front wall, an opening formed in the front wall, and a drive member reciprocally mounted in the injector. The adapter includes a first section and a second section that are rotatable relative to each other about a hinge axis generally perpendicular to a longitudinal axis of the adapter to an open position to allow loading of the syringe into the adapter from a position to the rear of the hinge axis. The first section and the second section are also preferably rotatable about the hinge axis to a closed position to form a syringe carrier to seat at least a portion of the syringe.

The present invention also provides an adapter for releasably mounting a syringe in a desired position relative to a powered injector. The adapter preferably includes a first section and a second section that are of generally the same in construction. The first section and the section are connectable to form a syringe carrier to seat at least a portion of the syringe. Preferably, the first section and the section also form a releasable mounting mechanism positioned to the rear of the syringe carrier to mount the adapter in a desired position relative to the front wall of the injector.

In another aspect, the present invention provides an adapter for releasably attaching a syringe to a front-loading powered injector including a releasable mounting mechanism positioned at a rear of the adapter to mount the adapter in a desired position relative to the front wall of the injector. The adapter also includes a syringe carrier section adapted to seat at least a portion of the syringe. The syringe carrier section is preferably open along the length of the top thereof to allow placement of the syringe therein from the top. A portion of the syringe carrier section abuts the transition region of the syringe in a manner that the force exerted by the syringe on the adapter is generally symmetrical about an axis of the adapter during an injection, thereby reducing any bending moment about the mounting mechanism. The syringe carrier section also includes an opening in a rear section thereof to allow the drive member of the injector to communicate forward force to the syringe plunger. The portion of the syringe carrier section abutting the transition region of the syringe can, for

example, include a first abutment surface positioned on a first lateral side of the syringe carrier section and a second abutment surface positioned on a second lateral side of the carrier section.

5 In another aspect, the present invention provides an adapter for releasably attaching a syringe to a front-loading powered injector including an intermediate section through which a push rod can pass to communicate force from the injector drive member to the plunger. The intermediate section has a releasable mounting mechanism positioned at a rear thereof to mount the adapter in a desired position relative to the front wall of the injector. The adapter also includes a syringe  
10 carrier section connected to the intermediate section. The syringe carrier section is adapted to seat at least a portion of the syringe and includes an opening in a rear section thereof to allow the drive member of the injector to communicate forward force to the plunger via the push rod.

The syringe carrier section is preferably open on a top thereof. A  
15 forward portion of the syringe carrier section abuts the transition region of the syringe during injection. The syringe carrier section is movable relative to the intermediate section to move the forward portion out of contact with the transition region of the syringe to enable removal of the syringe without retraction of the drive member. In one aspect, for example, the carrier section is connected to the intermediate section in a  
20 hinging manner.

In another aspect, an adapter system of the present invention includes an intermediate section having a releasable mounting mechanism positioned at a rear thereof as described above. The adapter system also includes a syringe carrier section connected to the intermediate section and adapted to seat at least a portion of the  
25 syringe. The syringe carrier section includes an opening in a rear section thereof to allow the drive member of the injector to communicate forward force to the plunger from the drive member and is preferably open on a top thereof. A forward portion of the syringe carrier section abuts the transition region of the syringe during an injection

procedure. The adapter system further includes a push rod used to communicate force from the injector drive member to the plunger. The pushrod has a forward section that is movable relative to a rearward section of the pushrod to allow movement of the transition region of the syringe out of contact with the forward portion of the syringe carrier section to enable removal of the syringe from the adapter without retraction of the drive member. The forward section of the push rod can, for example, be rotatable, hingeable or removable relative to the rearward section of the push rod.

Another adapter system of the present invention includes a releasable mounting mechanism positioned at a rear of the adapter to mount the adapter in a desired position relative to the front wall of the injector as described above and a pushrod to communicate force from the drive member to the plunger. The adapter system also includes a syringe carrier section adapted to seat at least a portion of the syringe. The syringe carrier section includes an opening in a rear section thereof to allow passage of the pushrod therethrough to contact to the plunger. The adapter system further includes a contact member preferably positioned in the syringe carrier section to contact the pushrod and prevent a fluid from the syringe from passing rearward of the sealing member. The contact member can, for example, include a wiper seal.

In another aspect, the present invention provides an adapter including a releasable mounting mechanism and a syringe carrier section adapted to seat at least a portion of the syringe. The syringe carrier section is preferably open on a top thereof to allow placement of the syringe therein or removal of the syringe therefrom from the top. The syringe carrier section includes at least one flexing retainer therein to place pressure on at least one side of the syringe to retain the syringe within the syringe carrier section. The syringe carrier section further includes an opening in a rear section thereof to allow the drive member of the injector to communicate forward force to the plunger.

In still a further aspect, the present invention provides an adapter for releasably attaching a syringe to a front-loading powered injector including a



releasable mounting mechanism as described above and a syringe carrier section adapted to seat at least a portion of the syringe. A forward abutment portion of the syringe carrier section abuts the transition region of the syringe. The syringe carrier section includes a biasing member to contact a rear surface of the syringe. The biasing member forces or biases the transition region of the syringe against the forward abutment portion for syringes of various lengths. The biasing member can, for example, be biased forward by a spring. The biasing member can also, for example, be biased forward by flexible member on a rear side thereof.

### **Brief Description of the Drawings**

Other aspects of the invention and their advantages will be discerned from the following detailed description when read in connection with the accompanying drawings, in which:

Figure 1A illustrates an embodiment of an injector system of the present invention for use in connection with an MRI procedure.

Figure 1B illustrates the injector system of Figure 1A in which the saline syringe and the adapter have been disassembled from the injector.

Figure 2A illustrates a perspective view of an embodiment of an adapter of the present invention in an open state for loading of a syringe therein.

Figure 2B illustrates a perspective view of the adapter of Figure 2A in an open state with a syringe loaded therein.

Figure 2C illustrates a perspective view of the adapter of Figure 2A in a closed state with a syringe loaded therein.

Figure 2D illustrates a plan view of the adapter of Figure 2A in a closed state with a syringe loaded therein.

Figure 2E illustrates a side, cross-sectional view of the adapter of Figure 2A in a closed state with a syringe loaded therein.

Figure 3A illustrates a perspective view of another embodiment of an adapter of the present invention in an open state for loading of a syringe therein.

5                Figure 3B illustrates a perspective view of the adapter of Figure 3A in an open state with a syringe loaded therein.

Figure 3C illustrates a perspective view of the adapter of Figure 3A in a closed state with a syringe loaded therein.

10              Figure 3D illustrates a plan view of the adapter of Figure 3A in a closed state with a syringe loaded therein.

Figure 3E illustrates a side, cross-sectional view of the adapter of Figure 3A in a closed state with a syringe loaded therein.

Figure 4A illustrates a perspective view of another embodiment of an adapter of the present invention with a syringe positioned to be inserted therein.

15              Figure 4B illustrates a perspective view of the adapter of Figure 4A with the syringe positioned therein.

Figure 4C illustrates a perspective view of the adapter of Figure 4A wherein a syringe retaining member is in a closed position.

20              Figure 4D illustrates a perspective view of an adapter including another embodiment of a syringe retaining member.

Figure 4E illustrates a perspective view of an adapter including another embodiment of a syringe retaining member.

Figure 4F illustrates a perspective view of an adapter including another embodiment of a syringe retaining member.

Figure 4G illustrates a perspective view of an adapter including another embodiment of a syringe retaining member.

5           Figure 4H illustrates a perspective view of an adapter including another embodiment of a syringe retaining member.

Figure 4I illustrates a perspective view of an adapter including another embodiment of a syringe retaining member.

10           Figure 5A illustrates a perspective view of an embodiment of an adapter including separate, generally identical sections in an unconnected state.

Figure 5B illustrates a perspective view of the adapter of Figure 5A in a connected state.

15           Figure 6A illustrates a perspective view of another embodiment of an adapter including generally identical sections that are hingingly attached via a side wall thereof.

Figure 6B illustrates a front view of the adapter of Figure 6A in an open state.

Figure 6C illustrates a perspective view of the adapter of Figure 6A in a closed state.

20           Figure 7A illustrates a perspective view of another embodiment of an adapter including generally identical sections that are attached at a front end thereof in an open state.

Figure 7B illustrates the adapter of Figure 7A with a syringe loaded therein.

Figure 7C illustrates the adapter of Figure 7A wherein the sections thereof have been hingingly closed for connection of the adapter to an injector.

5           Figure 8A illustrates a perspective view of a portion of an injector having at least one removable face plate for attachment of a syringe or an adapter thereto.

Figure 8B illustrates a perspective view of an embodiment of an adapter for use with the injector of Figure 8A.

10           Figure 8C illustrates a perspective view of another embodiment of an adapter for use with the injector of Figure 8A.

Figure 8D illustrates a perspective view of another embodiment of an adapter for use with the injector of Figure 8A.

Figure 8E illustrates a perspective view of another embodiment of an adapter for use with the injector of Figure 8A.

15           Figure 9A illustrates a perspective view of an embodiment of an adapter assembly or system in which a push rod performs the function of a syringe plunger extension rod.

Figure 9B illustrates a perspective view of the adapter assembly of Figure 9A in a disconnected state.

20           Figure 9C illustrates a cross-sectional view of the adapter assembly of Figure 9A.

Figure 10A illustrates a perspective view of an embodiment of an adapter assembly or system in which the carrier section hinges with respect to the

intermediate section to allow removal of the syringe after an injection procedure without retraction of the injector drive member.

Figure 10B illustrates an expanded perspective view of the forward portion of the carrier section.

5                Figure 10C illustrates a perspective view of an embodiment of a syringe retainer for use in the adapter system.

Figure 10D illustrates a front cross-sectional view of the syringe retainer of Figure 10C.

10              Figure 11 illustrates a perspective view of an adapter system in which the push rod hinges to allow removal of the syringe after an injection procedure without retraction of the injector drive member.

Figure 12A illustrates a perspective view of an embodiment of an adapter assembly in a disconnected state including a contact or sealing member for removing injection fluids from a push rod.

15              Figure 12B illustrates a perspective view of the adapter assembly similar to the adapter assembly of Figure 12A in a connected state.

Figure 13A illustrates a perspective view of an embodiment of an adapter assembly including a biasing member to bias the syringe forward within the carrier section.

20              Figure 13B illustrates a perspective view of the adapter assembly of Figure 13A in a disconnected state.

Figure 14 illustrates a perspective view of another embodiment of a biasing member to bias the syringe forward within the carrier section.

### Detailed Description of the Invention

An embodiment of a front-loading injector system 5 of the present invention is illustrated in Figures 1A and 1B. Injector system 5 is particularly adapted for use in MRI procedures and includes a powered injector 10, a syringe 20 for injection  
5 of saline solution and an adapter 100. An example of an injector 10 suitable for use in the present invention is the SPECTRIS® injector available from Medrad, Inc. of Indianola, Pennsylvania. However, the present invention may be used in connection with other fluid delivery systems, including injectors and infusion pumps for computed tomography, ultrasound and angiographic procedures. As best illustrated in Figure 1B,  
10 injector housing 30 of injector 10 preferably includes a first drive member or piston 40 therein which cooperates with a syringe plunger 50 in saline syringe 20 to inject a saline solution from the interior of syringe 20 into a patient.

As shown in Figures 1B, injector 10 also includes a second drive member or piston 40' that cooperates with an adapter 100 and a syringe plunger  
15 extension rod 220 of a syringe 200 (see, for example, Figure 2A) containing a fluid such as a contrast medium to inject the fluid from the interior of syringe 200 into a patient.

As used herein to describe injection system 5 and other embodiments of the present invention, the terms "axial" or "axially" refer generally to, for example, an axis A around which adapter 100 is preferably formed (although not necessarily  
20 symmetrically therearound) or an axis B around which saline syringe 20 is formed (although not necessarily symmetrically therearound). The terms "proximal" or "rearward" refer generally to an axial or a longitudinal direction toward the end of injector housing 30 opposite the end to which syringe 20 and adapter 100 are mounted. The terms "distal" or "forward" refer generally to an axial or a longitudinal direction  
25 toward a syringe tip of syringe 20 or syringe 200. The term "radial" refers generally to a direction normal to an axis such as axis A or axis B.

Saline syringe 20 and adapter 100 are preferably removably connected to injector 10 as described in U.S. Patent No. 5,383,858. In that regard, front-loading

injector 10 preferably includes a front wall 60 having a first opening 62 formed therein. Piston 40 is reciprocally mounted within injector 10 and is extendible through opening 62. Piston 40 preferably includes a piston flange or head 44. Receiving slots 66a and 66b, are preferably positioned opposite one another around opening 62.

5 Receiving flanges 68a and 68b are preferably positioned opposite one another and between receiving slots 66a and 66b and extend inwardly into opening 62.

The rearward end of saline syringe 20 preferably includes a readily releasable mounting mechanism such as a pair of mounting flanges 22a and 22b for mounting saline syringe 20 in a desired position relative to the front wall 60 of injector 10. Flange 22b is not shown but is generally identical to flange 22a and positioned opposite flange 22a. Mounting flanges 22a and 22b may include indicating means, such as detent(s), bar code(s), protrusion(s) or notch(es) 24a, which provide information to the injector 10, for example, about the type of saline syringe 20 being used. Correspondingly, injector 10 preferably includes any suitable means (not

10 shown) for reading information from notch(es) 24a.

15

To attach syringe 20 to injector 10, the rearward end of syringe 20 is inserted into injector opening 62 such that mounting flanges 22a and 22b are inserted into receiving slots 66a and 66b, respectively. If, at this time, plunger 40 is not positioned at the rearward end of syringe 20 such that a piston flange 44 can engage capture members 54 (as described in U.S. Patent No. 5,383,858), piston 40 may be advanced forward by the operation of injector 10 until piston flange 44 is in position to be received by capture members 54.

20

Once mounting flanges 22a and 22b are inserted into receiving slots 66a and 66b, respectively, and piston 40 is in position to be received by capture members 54, the operator preferably rotates syringe 20 approximately 90 degrees such that mounting flanges 22a and 22b move behind and are engaged by receiving flanges 68a and 68b, respectively, and piston flange 44 rotates into position to be retained by, for example, L-shaped capture members 54. Injector 10 may include a stop mechanism (not shown), for example, extending from at least one of the retaining

25

slots 68a and 68b, to prevent rotation of syringe 20 more than 90 degrees. Tactile, visual or audible feedback can be provided to the operator via, for example, cooperating members on syringe 20 and injector 10 to inform the operator that secure connection has been achieved. After securely attaching syringe 20 to injector 10, advancing piston 40 in a forward direction will apply a motive force to plunger 50 to advance plunger 50 forward within syringe 20, thereby forcing saline solution in syringe 20 out of syringe neck 26 into the fluid path to the patient. Retracting piston 40 in a rearward direction will cause plunger 50 to move rearward in syringe 20, thereby drawing fluid into syringe 20.

Adapter 100 is preferably attached to injector 10 in a similar manner as described above for attachment of syringe 20 to injector 10. In that regard, a rearward portion or section of adapter 100 preferably includes a readily releasable mounting mechanism such as a pair of mounting flanges 102a and 102b (see Figure 2A) for mounting adapter 100 in a desired position relative to the front wall 60 of injector 10. Mounting flanges 102a and 102b may include indicating means, such as detents or notches 104a and 104b, which provide information to injector 10 about the type of adapter and/or syringe being used. Correspondingly, injector 10 preferably includes any suitable means (not shown) for reading information from notches 104a and 104b.

To attach adapter 100 to injector 10, the rearward end of adapter 100 is inserted into injector opening 62' such that mounting flanges 102a and 102b are inserted into receiving slots 66a' and 66b', respectively. Once mounting flanges 102a and 102b are inserted into receiving slots 66a' and 66b', respectively, the operator preferably rotates adapter 100 or adapter 100/syringe 200 combination approximately 90 degrees such that mounting flanges 102a and 102b move behind and are engaged by receiving flanges 68a' and 68b', respectively. As described above, a stop mechanism (not shown) may, for example, extend from at least one of the retaining slots 68a' and 68b', to prevent rotation of adapter 100 more than 90 degrees. Once again, tactile, visual or audible feedback can be provided to the operator via, for example, cooperating members on adapter 100 and injector 10 to inform the operator that secure connection has been achieved. A drip flange 106 can, for example, be



formed on a rearward portion of adapter 100 to, among other things, assist in forming a secure connection. Drip flange 106 may, for example, include a raised member or detent 108 (see, for example, Figure 2A) that mates with a recess (not shown) in the face of opening 62' to provide audible and/or tactile feedback to the operator upon proper alignment/connection of adapter 100 to injector 10.

After securely attaching adapter 100 to injector 10, advancing piston 40' in a forward direction will apply a motive force to a plunger extension 220 of syringe 200 to advance syringe plunger 225 (see Figure 2E) forward within syringe barrel 210, thereby forcing contrast medium in syringe 200 out of syringe neck 250 into the fluid path to the patient.

Adapter 100 is illustrated in further detail in Figures 2A through 2E. In the embodiment of adapter 100, a "break" action is used to load syringe 200 into a carrier 110 of adapter 100. In that regard, carrier 110 includes a first portion or section 120 and a second portion or section 130. First portion 120 is hingingly attached to second portion 130 via support arms 132a and 132b, each of which includes a passage 134 therein. First portion 120 includes generally cylindrical tabs 122 on each side thereof that snap into passages 134 to hingingly or rotatably attach first portion 120 to second portion 130 about an axis C (see, for example, Figure 2D) preferably oriented generally perpendicular to longitudinal axis A of adapter 100.

Figure 2A illustrates adapter 100 in an open state and ready to receive syringe 200 from a position to the rear of the hinging mechanism. In this embodiment, syringe 200 comprises generally cylindrical body or barrel 210 in which a fluid such as contrast medium, saline or therapeutic agent is contained. Preferably, the fluid medium is "prefilled" into syringe 200 before loading of syringe 200 in adapter 100. Syringe 200 can, for example, be prefilled by the manufacturer or manually filled remote from the injector. Syringe 200 further includes plunger 225 slideably disposed within barrel 210 that is similar in operation to plunger 50 of saline syringe 20. Plunger 225 of syringe 200 is operatively connected to plunger extension rod 220 by, for example, a threaded connection. Syringe 200 further includes a flange 230 at a rearward

end of barrel 210. At a forward end of barrel 210, syringe 200 includes, for example, a generally frusto-conical transition or cone region 240 that connects barrel 210 to a tapered neck 250 from which contrast medium is injected. Tapered neck 250 can include, for example, a luer connection at the end thereof for connection to a fluid path  
5 (for example, flexible tubing) as known in the art.

In many cases, syringes 200 for use, for example, in an MRI procedure are prefilled with contrast medium by the manufacturer. Many such syringes 200 are designed for manual injection into a patient wherein an operator manually advances plunger rod 220 (and thereby plunger 225 within syringe 200) forward by applying  
10 pressure to a rearward end 228 of plunger extension rod 220. Syringe barrel 210 and flange 230 may, for example, be fabricated from glass or plastic. Plunger extension rod 220 is typically fabricated from a plastic material.

As illustrated in Figures 2A and 2B, syringe 200 is loaded into carrier 110 by positioning syringe 200 in general alignment with a generally cylindrical  
15 passage 140 formed in a forward end of first portion 120 of adapter 100. Syringe 200 is slid forward within passage 140 until syringe flange 230 abuts a shoulder 150 (see Figure 2E) that extends radially inwardly within first portion 120. Shoulder 150 cooperates with syringe flange 230 to hold syringe 200 within adapter 100 and to provide resistance to the forward force applied to plunger extension rod 220 by  
20 piston 40' during an injection procedure.

As best illustrated in Figure 2E, shoulder 150 preferably includes a rearward extending ridge portion 154 on the inner radius thereof so that ridge  
portion 154 contacts flange 230 only near or in the vicinity of the transition of syringe flange 230 into syringe barrel 210. Contact of ridge portion 154 with syringe flange 230  
25 near barrel 210 assists in minimizing the forces placed upon cantilevered flange 230. For example, in the case of glass syringes 200, flange 230 may fail (break) if contact with shoulder 150 is made at or near the outer end of syringe flange 230, whereas the shorter lever arm resulting from contact with syringe flange 230 at or near the inner radius thereof will reduce the force on syringe flange 230 and prevent failure. Ridge

portion 154 can be made of a resilient or compliant material such as an elastomeric material that can be different from the material of the remainder of adapter 100 to further reduce the likelihood of failure.

5 In the case of a prefilled syringe 200, there is no need for the operator to retract the plunger of syringe 200 to load syringe 200 with contrast medium. Therefore, there is usually no need for a follower mechanism in the adapter of the present invention to attach to plunger extension rod 220 to enable retraction of plunger 225 as described in connection with the adapter of U.S. Patent No. 5,520,653. Piston 40' can simply be advanced to abut rearward surface 228 of plunger extension rod 220. Any further  
10 forward motion of piston 40' will result in advancement of the plunger of syringe 200 and pressurization of the contrast medium with syringe 200. Elimination of a carrier mechanism for the plunger extension simplifies and reduces the cost of manufacture of the adapters of the present invention as compared, for example, to the adapter of U.S. Patent No. 5,520,653. Nevertheless, the adapters of the present invention may readily  
15 be configured with follower mechanisms that connect to plunger extension rods 220 to allow plunger retraction.

Figures 2C through 2E illustrate syringe 200 within adapter 100 with first portion 120 and second portion 130 in a closed position. Adapter 100 preferably includes a mechanism to assist in maintaining first portion 120 and second portion 130  
20 in a closed position during operation of injector 100. Second portion 130 may, for example, include a latch tab 136 having an abutment shoulder 138 that cooperates with a recess 128 in a rearward end of first portion 120 to create a snap latching mechanism. Many other closing mechanisms can be used to maintain first portion 120 and second portion 130 in a closed position, as clear to one skilled in the art.

25 Closed carrier 110 created by first portion 120 and second portion 130 also functions to limit the motion of plunger extension rod 220 out of alignment with axis A. This prevents plunger extension rod 220 from, for example, slipping out of contact with piston 40', prevents deforming of plunger extension rod 220 and prevents eccentric loading of plunger extension rod 220. Deflection, eccentric loading or

deforming of plunger extension rod 220 may, for example, cause leaking of fluid to the rear of plunger 225 or breaking of syringe 200. Radially inward projecting guide(s) can be formed in one or both of first portion 120 and second portion 130 to maintain even tighter tolerances. A section of either or both of first portion 120 and second  
5 portion 130 can be "cut away" to form a window for viewing of syringe extension rod 220. Likewise, a portion or the entirety of either or both first portion 120 and second portion 130 can be transparent.

Preferably, one or both of first portion 120 and second portion 130 includes an abutment member to prevent rotation of syringe 200 within carrier 110. It  
10 is, for example, desirable to prevent rotation of syringe 200 after connection thereof to fluid path tubing. Prevention of syringe rotation can also maintain syringe 200 in proper orientation for viewing, for example, volume gradations on syringe 200. One or more sides of first portion 120 can, for example, have a flattened profile to conform to flattened section(s) 234 of syringe flange 230 to prevent rotation of syringe 200.

15 As illustrated in Figures 2B through 2E, a substantial portion of syringe 200 extends forward through passage 140 so that syringe barrel 210 is plainly visible to the operator. Such visibility, for example, facilitates reading of wording printed on the syringe as well as visual determination of the volume of contrast remaining in syringe 200. Visibility of syringe 220 also allows the operator to more  
20 readily determine whether air is present in syringe 200 before commencing an injection procedure. Moreover, operators typically like to see plunger 225 in motion to provide reassurance that the injection is proceeding. Extension of a portion of syringe 200 beyond carrier 110 also facilitates grasping of syringe 200 by the operator to, for example, connect or disconnect a fluid path to syringe neck 250.

25 As illustrated, for example, in Figures 2A through 2C, axis C of rotation of the hinge mechanism (that is, the axis passing through the radial center of generally cylindrical tabs 122 in the embodiment of Figure 2A) is preferably positioned such that the force experienced by carrier 110 during forward advancement of piston 40' tends to force or maintain carrier 100 in a closed position. The axis of

cylindrical tabs 122 is preferably, for example, positioned above the center line or longitudinal axis of carrier 110 such that a forward force exerted on shoulder 150 tends to produce a torque that maintains first portion 120 in a latched, closed position relative to second portion 130.

5                Once an injection procedure is completed, the operator can grasp the adapter or adapter/syringe combination and rotate it 90 degrees back to the pre-installation orientation, thereby, disengaging mounting flanges 102a and 102b from behind receiving flanges 68a and 68b, respectively. The adapter/syringe combination is then removable from the injector 10.

10              Retaining syringe 200 within carrier 110 by abutment with shoulder 150, allows accommodation of many different designs of syringe 200 by carrier 110. Adapter 100 is thus usable with a wide variety of currently available syringes 200.

              Figures 3A through 3E illustrate another embodiment of an  
15              adapter 300 for use with a syringe 200. The rearward portion of adapter 300 is essentially identical to that of adapter 100 and is removably attached to injector 10 as described above. Unlike adapter 100, which holds syringe 200 within adapter 100 and provides resistance to the forward force applied to plunger extension rod 220 by abutment of syringe flange 230, syringe 200 is held within adapter 300 and resistance  
20              provided to the forward force applied to plunger extension rod 220 by abutting a forward facing surface of forward transition region 240 of syringe 200 rather than by abutting or retaining syringe flange 230.

              Like adapter 200, a hinging or "break" action is used to load syringe 200 into a carrier 310 of adapter 300. In that regard, carrier 310 includes a first portion 320  
25              and a second portion 330. First portion 320 is hingingly or rotatably attached to second portion 330 via support arms 332a and 332b, each of which includes a passage 334 therein. First portion 320 includes generally cylindrical tabs 322 on each side thereof that reside in passages 334 to hingingly attach first portion 320 to second portion 330.

First portion 320 rotates about an axis C that runs generally through the radial centers of tabs 322 and is generally perpendicular to the longitudinal axis A of adapter 300.

Figure 3A illustrates adapter 300 in an open state and ready to receive syringe 200. As illustrated in Figures 3A and 3B, syringe 200 is loaded into carrier 310 by positioning syringe 200 in general alignment with a generally cylindrical passage 340 formed in first portion 320 of adapter 300. Syringe 200 is slid forward within passage 340 until a forward facing surface of syringe 200 abuts a retention shoulder 350 (see Figure 3E) that extends radially inward at a forward end of a forward portion 342 of first portion 310. Shoulder 350 cooperates with forward syringe transition region or cone 240 to hold syringe 200 within adapter 300 and to provide resistance to the forward force applied to plunger extension rod 220 by piston 40' during an injection procedure. As illustrated in detail D of Figure 3E, shoulder 350 preferably contacts syringe 200 only near to or in the vicinity of the transition from the sidewall of barrel 110 to transition region 240 to take advantage of increased structural strength in this region. The contact area of shoulder 350 can be made of a resilient or compliant material (for example, an elastomeric material) to absorb energy and reduce the likelihood of breaking syringe 200.

Syringe 200 may be inserted in adapter 300 before connection of adapter 300 to injector 10. Alternatively, syringe 200 may be loaded into adapter 300 while adapter 300 is mounted on injector 10. Indeed, adapter 300 may remain mounted on injector 10 through many different injection procedures with different syringes.

Forward portion 342 preferably includes one or more open areas or windows 344 so that syringe barrel 210 is plainly visible to the operator. All or a portion of forward portion 342 can also be transparent to further facilitate viewing of syringe barrel 210. Open areas 344 also facilitate grasping of syringe 200 by the operator to, for example, connect or disconnect a fluid path to syringe neck 250. As described in connection with adapter 100, one or both of first portion 320 and second portion 330 may include an abutment member to prevent rotation of syringe 200 within carrier 310. For example, the side(s) of first portion 320 can have a flattened

profile to conform to flattened section(s) 234 of syringe flange 230 to prevent rotation of syringe 200. The cooperation of such a flattened profile of carrier 310 and section 234 can, for example, be used to ensure a desired orientation of syringe barrel 210 with open areas 344. For example, two open areas 344 can be provided  
5 generally opposing each other (that is, positioned approximately 180° apart on forward portion 342). The cooperation of a flattened profile of carrier 310 and flattened syringe flange section 234 in this embodiment preferably allows mounting of syringe 200 in carrier 310 in only two axially rotated orientations, 180° apart.

One or a plurality of inward projecting guide 360 can be formed in one  
10 or both of first portion 120 and second portion 130 to maintain tight tolerances to prevent deflection of plunger extension rod 330 as discussed above. A plurality (for example, three) guides 360 can be used about first portion 120 and/or second portion 130 to limit or prevent deflection in any direction.

Open areas (not shown) can also be provided on carrier 310 in the area  
15 where syringe flange 230 resides to accommodate large (in a radial direction) or irregularly shaped syringe flanges 230. Such open areas preferably extend longitudinally to accommodate syringes of different length from a forward end thereof to the syringe flange thereof. In general, the adapters of the present invention preferably provide adequate capacity to accommodate syringes of widely varying  
20 length, diameter etc.

An important function of an injector is to monitor and report the actual volume of fluid available for delivery within a syringe. This function, for example, enables rapid decision on whether enough fluid is present to proceed with an imaging procedure or whether additional volume should be loaded. Monitoring the cumulative  
25 volume of fluid delivered to a patient is also desirable for certain applications where a recommended per-patient dosage volume should not be exceeded. Fluid volumes delivered by injectors are typically displayed in 1.0 ml increments and are tracked by the injector with finer resolution than is displayed. Injectors also preferably detect

and differentiate among different types and sizes of syringes so that accurate display and delivery of fluid volume is provided.

To achieve such fluid volume management specifications, an adapter or syringe must be installed on an injector, be oriented in a known manner and provide the control system with identification information. Identification can be provided by coded features on the adapter or syringe that are detected by sensors on the injector so that each adapter and/or syringe is known by its code and fluid volume parameters specific to that syringe are implemented. When syringes 200 that are intended for non-powered, hand injection are installed on power injector 10, adequate identification is still preferably provided. This result can be achieved by first placing syringe 200 in adapter 100 or 300 which facilitates mounting on injector 10 and possesses its own unique code as encoded by, for example, the positions of notches 104a and 104b. In this manner, many different types and sizes of syringes 200 and/or adapters can be accommodated.

Syringes 200 are frequently of similar geometry such that more than one type of syringe 200 can be carried by the same adapter, giving rise to the potential of incorrect identification and possible fluid delivery error by injector 10. However, it is desirable to minimize the number of adapters required to accommodate all the hand syringes intended for a specific injector leading to a need to install more than one volume of syringe per adapter, provided individual functionality can be achieved. Syringes that share the same functional internal diameter, but have different lengths of travel can be treated as equivalent if both are referenced to the adapter, and consequently to injector 10, by a front-most surface thereof (that is, cone or transition region 240). If syringes 200 sharing the same functional diameter but having different lengths are referenced to the adapter injector 10 by rear flange 230 only, injector 10 will not be able to determine where the front of a syringe 200 is and cannot accurately determine/report the volume of contrast medium available. Mounting hand syringes 200 by rear flange 230 as in the embodiment of adapter 100 thus preferably requires one adapter per syringe diameter and length combination, which results in a larger number of adapters than would be required using a front mounting as in the



embodiment of adapter 300. Multiple adapter combinations decrease the ease of use for an operator and expand the logic and sensing capacity required of injector 10. An optimum approach would be to use a single adapter that accommodates all hand syringes targeted for a particular injector. To approach this goal, it is preferable to retain/about a front end of syringes 200 as described above in connection with adapter 300 so that injector 10 can determine the position of the front end of syringe 200. Loading of a front end of syringe 200 is also preferred to take advantage of an area of increased syringe strength to prevent syringe failure.

Figures 4A through 4C illustrate another embodiment of an adapter 400 for use with syringe 200. The rearward portion of adapter 400 is essentially identical to that of adapters 100 and 300 and is removably attached to injector 10 as described above. Unlike adapters 100 and 300, which incorporate a hinging action to enclose syringe 200, syringe 400 includes an open carrier 410. Like adapter 300, however, syringe 200 is held within adapter 400 and resistance provided to the forward force applied to plunger extension rod 220 by abutting forward transition or cone region 240 of syringe 200.

As illustrated in Figures 4A and 4B, syringe 200 is simply loaded into adapter 400 by dropping syringe 200 therein from above. Adapter 400 preferably includes a first, rearward portion 420 that seats/supports syringe flange 230. Preferably, first portion 420 has generally flat side walls 422 that cooperate with generally flat sections 234 on syringe flange 230 to restrict or substantially prevent rotation of syringe 200 within adapter 400. Side walls 422 of first portion 420 preferably extend upward past the generally common axis of syringe 200 and adapter 400 to assist in supporting syringe 200.

Adapter 400 further includes a second, forward portion 430 that seats/supports syringe barrel 210. Second portion 430 preferably includes a radially inward extending abutment shoulder 450 that abuts cone or transition region 340 of syringe 200 to retain syringe 200 within adapter 400 and provide resistance to the forward force applied to plunger extension rod 220 by piston 40'. Although second

portion 430, including abutment shoulder 450, are open on the top thereof, the generally cylindrical wall of second portion 430 and abutment shoulder 450 can extend upward past the generally common axis of syringe 200 and adapter 400 to support resistance of the forward force applied to plunger extension rod 220 and to prevent deflection of syringe 200 out of alignment with the shared axis of syringe 200 and adapter 400.

Through abutment shoulder 450, adapter 400 provides the benefits of forward abutment of syringe 200 discussed above. Moreover, because adapter 400 is open along its entire axial length, insertion and removal of syringe 200 is facilitated. For example, syringe 200 can even be easily removed from adapter 400 while still connected to a fluid path (not shown in Figures 4A through 4C). In some cases, however, it may be desirable to form abutment shoulder 450 to contact a forward facing surface of transition region 240 around its entire circumference to provide additional stability. In that case, the forwardmost end of carrier 410 would be closed and any fluid path would preferably be disconnected before removing syringe 200 from adapter carrier 410.

Adapter 400 preferably further includes at least one retaining member 460 to assist in retaining and/or stabilizing syringe 200 in proper alignment therein. Retaining member 460 is slideably retained in a generally cylindrically shaped passage 470 in carrier 410. Retaining member 460 is illustrated in an open or disengaged position in Figures 4A and 4B. To close or engage retaining member 450 to retain syringe 200, the operator can supply force to collar tab 462 to rotate retaining member 450 within passage 470 to a closed position as illustrated in Figure 4C. In another embodiment illustrated in Figure 4D, rotatable retaining member 460 can be split into two portions 460' and 460'' that rotate to meet in the middle.

Other retaining/stabilizing members or mechanisms for retaining/stabilizing syringe 200 are illustrated in Figures 4E through 4I. In Figure 4E a sliding retaining member 460a is positioned on/around second, forward portion 430 of carrier 410. Retaining member 460a is preferably positioned at a forwardmost

position on second portion 430 when syringe is loaded into carrier 410 to facilitate loading. Retaining member 460a is retained on second portion 430 by abutment with a forward shoulder 480 and rearward shoulder 482. After seating of syringe 200, retaining member 450a can preferably be slid to any desired position on second  
5 portion. Positioning retaining member 460A at a rearwardmost position on second portion 430 may maximize stability. Retaining member 460a preferably conforms closely to the shape of syringe barrel 210 to maximize stability. Retaining member 460a may include an open section 462a on the top thereof to facilitate removal of syringe 200 from carrier 410 without disconnection of an attached fluid  
10 path element.

Figure 4F illustrates another embodiment of a slideable retaining member 460b. Retaining member 460b is slideably retained upon first portion 420 of carrier 410 between drip flange 106 and shoulder 484. To facilitate loading of syringe 200, retaining member 450b may be positioned near drip flange 106. After  
15 loading of syringe 200, retaining member 460b can be slid to a desired position. An opening (not shown) in retaining member 460b can be formed to facilitate removal of syringe 200 from carrier 410 without disconnection of an attached fluid path element.

In the embodiment of Figure 4G, a retaining mechanism includes two cantilevered retaining member 460c' and 460c'' that snap around syringe barrel 210  
20 upon loading of syringe 200 in carrier 410.

A plurality of retaining/stabilizing members as described above can be provided along the length of carrier 410 to assist in retaining/stabilizing syringe barrel 210 and plunger extension rod 420 in proper position within carrier 410. The opening and closing of such retaining members can be operated individually or  
25 collectively, for example, via a common tab.

Alternatively, a retaining/stabilizing member can be increased in axially length to increase stability. For example, Figure 4H illustrates a retaining member 460d hingingly attached to first portion 420 that extends along the entire

length of first portion 420 and partially along the length of second portion 430 when closed. A similar retaining member can alternatively or additionally be hingingly attached to second portion 430. As an alternative to a hinging motion, such retaining member can rotate in a passage or slot formed in carrier 410.

5           An example of a widened, rotating retaining member 460e is illustrated in Figure 4I. Retaining member 460e is rotatably attached (about the longitudinal axis of carrier 410) to second portion 430 and extends generally along the entire length of second portion 430. After syringe 200 is top loaded into carrier 420, retaining member 460e can be rotated to form a cover over second section 430. Retaining members 460d and 460e can be transparent or formed with cut away sections to  
10           enhance the visibility of syringe 200.

          In some cases it may be desirable to manufacture the adapter of the present invention to be disposable after one or more uses. Such disposable adapters are preferably manufactured in an inexpensive manner. In Figures 5A and 5B, an  
15           embodiment of a preferably disposable adapter 500 is illustrated. Adapter 500 preferably includes two generally identical members 510a and 510b. Fabricating adapter 500 from two generally identical members 510a and 510b may substantially reduce manufacturing costs. Members 510a and 510b may for example be fabricated to "snap" together (for example, via extension members 520 and cooperating catch  
20           members 530) and provide visible, audible and/or tactile feedback to indicated proper connection.

          In use, syringe 200 is preferably seated in one of members 510a or 510b. The other portion is then, for example, snapped into place to encompass syringe 200 within the resultant adapter 500 (see Figure 5B). When assembled,  
25           portions 510a and 510b form a rearward section that houses plunger extension rod 220 and a forward portion that houses syringe barrel 210. At or near a rearward end of adapter 500, members 510a and 510b preferably form a connecting section including first mounting flange 502a and a second mounting flange 502b adapted to removably attach adapter 500 to injector 10 as described above. As illustrated in Figure 5b,

adapter 500 encompasses both syringe barrel 210 and plunger extension rod 220 to retain/stabilize syringe 200. The side walls of adapter 500 can be formed with a generally flat or flattened profile to interact with/about generally flat section(s) 234 of syringe flange 230 to prevent rotation of syringe 200 about its axis within adapter 200.

5 To facilitate viewing of either plunger extension rod 220 or barrel 210, adapter 500 can be formed with cut out window sections 540. Moreover, any portion or all of adapter 500 can be transparent.

Adapter 500 provides resistance to the forward force applied to plunger extension rod 220 by abutment of syringe transition region 240 with a radially inward  
10 extending shoulder section 550. The advantages of providing such resistance/retention by abutment of syringe transition region 240 discussed above in connection with other embodiments of adapter of the present invention are also provided by adapter 500.

Another embodiment of an adapter 500' is illustrated in Figures 6A  
15 through 6C. Adapter 500' is generally identical to adapter 500 except that adapter 500' is formed from the connection of a first portion 510a' and a second portion 510b' that are initially hinged together, for example by a notched plastic hinge 512 as known in the art) as best illustrated in Figures 6A and 6B.

Figures 7A through 7C illustrate another embodiment of an  
20 adapter 600 of the present invention. Adapter 600 includes a first member 610a and a second member 610b that are attached via a forward hinge mechanism. The hinge mechanism preferably includes a generally cylindrical member 612 on an extending member 614 of one of first member 610a and second 610b that is rotatably seatable in a passage 616 formed in an extending member of the other of first member 610a and  
25 second member 610b. Extending members 614 preferably extend forward on each side of a forward end of each of first member 610a and second member 610b as illustrated in Figure 7A.

First member 610a includes first retaining flange 102a as described above, while second member 610b includes second retaining flange 102b that operate to connect adapter 600 to injector 10 as described above. First member 610a preferably includes one half of a drip flange 106a, while second member 610b preferably includes the other half of a drip flange 106b.

To seat syringe 200 within adapter 600, first member 610a and second member 610b are preferably rotated about an axis C' of hinging mechanism (which axis is generally perpendicular to the orientation of longitudinal axis A' of adapter 600) to an open position as illustrated in Figures 6A and 6B. Barrel 210 of syringe 200 is then passed through an opening 640 formed at the forward end of adapter 600. Syringe 200 is advanced forward through passage 640 until flange 230 abuts radially inward extending shoulder 650 to retain syringe 200 within adapter 600 and provide resistance to forward force exerted upon plunger extension rod 220 by piston 40'. First member 610a and second member 610b are then rotated to a closed position. As discussed above in connection with adapter 100, shoulder 650 may include a raised (rearward extending) abutment ridge 660 to ensure that contact is made with syringe flange 230 near the point where syringe flange 230 is connected to barrel 210. Forward force exerted upon shoulder 650 assists in maintaining adapter 600 in a closed position as illustrated in Figure 6C.

Like adapters 500 and 500', first member 610a and second member 610b can be formed to be generally identical. Fabrications costs of adapter 600 can thereby be reduced.

In all of the embodiments discussed above, the adapter is attached to injector 10 via mounting flanges on a rearward portion or section of the adapter. There are, however, alternative manners in which an adapter of the present invention may be attached to injector 10. As illustrated in Figure 8A, for example, an injector may include a front wall 700 to which at least one removable face plate 705 is attached. The injector of Figure 8A is designed for use in an MRI procedure and includes a first, contrast syringe 710 attached to a first face plate 705 and a second,

saline syringe 720 attached to a second face plate 705'. In the embodiment of Figure 8A, face plate 705 is rotated upward to be detached from the injector.

Figure 8B illustrates an embodiment of an adapter 800 including a carrier 810 formed integrally or attached to a face plate 805 suitable for attachment to the injector of Figure 8A. Syringe 200 can be "breach" loaded into carrier 810 by first tilting syringe 200 and advancing barrel 210 of syringe 200 in a forward direction through passage 840 formed in a forward most position of an enclosed forward section 830 of carrier 810. Syringe 200 is advanced until syringe cone region 240 abuts radially inwardly extending shoulder 850 that defines passage 840. Face plate 805 includes a passage 808 therein through which a piston (not shown) of the injector of Figure 8A can pass to cooperate with plunger extension rod 220.

Figure 8C illustrates an embodiment of an adapter 900 that includes a carrier 910 attached to a removable face plate 905. In this embodiment, syringe 200 is advanced through passage 908 in face plate 902. Syringe 200 is advanced forward until cone region 240 abuts radially inward extending shoulder 950 of carrier 910.

Figure 8D illustrates an embodiment of an adapter 1000 that includes a carrier 1010 attached to a removable face plate 1005. Syringe 200 is loaded into carrier 1010 from the top by dropping syringe 200 into carrier 1010. When seated in carrier 1010, syringe 200 abuts radially inward extending shoulder 1050 of carrier 1010. The top of carrier 1010 is maintained in an open state over the length of carrier 1010 to facilitate removal of syringe 200 even when connected to a fluid path element.

Figure 8E illustrates an embodiment of an adapter 1100 that includes a carrier 1110 attached to a removable face plate 1105. Like adapter 1010, syringe 200 is loaded into carrier 1110 from the top by dropping syringe 200 into carrier 1110. When seated in carrier 1110, syringe transition region 240 abuts radially inward extending shoulder 1150 of carrier 1110. Carrier 1110 includes a hinging cover section 1160 that can be rotated to a closed position to form a cover/retainer over at

least a portion of syringe barrel 210 to assist in retaining/stabilizing syringe 200. Shoulder 1050 can be rotatable to capture syringe 200 and support it.

Figures 9A through 9C illustrate an embodiment of an adapter system 1200 including a carrier section 1210, an intermediate section 1220, and a rearwardmost connecting section 1230. Adapter 1200 further includes a push rod 1240. Syringe 200 is seated in carrier section 1210 by dropping syringe 200 into carrier section 1210. Cone region 240 abuts shoulder 1250 of carrier section 1210. The plunger extension rod (not shown) has been removed from connection with syringe plunger 225. In many cases, such plunger extension rods are connected to plunger 225 via threading on the forward end of the plunger extension rod.

Push rod 1240 extends through intermediate section 1220 to cooperate with plunger 225 to apply force to plunger 225. In the case of a prefilled syringe, there is typically no need to retract plunger 225 within syringe 200. In such cases, there is no need to establish an engaging connection (threaded or otherwise) between a forward end of push rod 1240 and plunger 225 to resist the force of and couple plunger 225 to push rod 1240 during a retracting motion of push rod 1240. This greatly simplifies the construction and operation of push rod 1240 and the injector.

In operation of adapter system 1200, push rod 1240 makes a connection with a piston (not shown in Figures 9A through 9C; see Figure 1A) of the injector through a connective coupling 1242 on a rearward end of push rod 1240. Connecting section 1230 is removably attached to the injector via the cooperation of mounting flanged 1202a and 1202b and drip flange 1206 with the injector as described above. Syringe 200 can be top loaded into carriage section 1210 either before or after connection of adapter 1205 to the injector via connecting section 1230. Push rod 1240 is advanced forward through intermediate section 1220 by the injector piston until forward end 1244 pilots into syringe plunger 225 to abut a rearward facing wall section within plunger 225. Once again, no secured connection to resist a rearward motion need be effected between push rod forward end 1244 and plunger 225 in, for example, the case of a prefilled syringe or in any other case that



retraction of plunger 225 within syringe 200 will not be required. Push rod forward end 1244 is preferably of generally the shape of the rearward facing interior of plunger 225. In this manner, push rod forward end 1244 provides support to plunger 225 to maintain the shape of plunger 225 during use of syringe 200. In many cases, plunger 225 will be fabricated predominantly from an elastomeric cover material. If the side walls of plunger 225 do not make adequate sealing contact with the interior side wall of syringe barrel 210, leakage of contrast to the rear of plunger 225 can occur during advancement of plunger 225.

Figures 10A through 10D illustrate another embodiment of an adapter system 1300 of the present invention. Similar to adapter system 1200, adapter system 1300 includes a carrier section 1310, an intermediate section 1320, and a rearwardmost connecting section 1330. Adapter system 1300 further includes or operates with a push rod 1340. Syringe 200 is seated in open carrier section 1310 by placing syringe 200 into carrier section 1310 from above (and can be seated therein or removed therefrom without removal or any attached tubing). Forward transition or cone region 240 abuts a shoulder portion 1350 of carrier section 1310.

To minimize fabrication costs of adapter system 1300, it is desirable that the option of using, for example, less expensive, lower-strength polymeric materials be available. Because the top portion of carrier section 1310 and shoulder 1350 are open for ease of removal of syringe 200, asymmetrical loading of connecting section 1330 can occur if cone region 240 of syringe 200 contacts a bottom portion of shoulder portion 1350 (as in the case of shoulder 1240, for example) during advancement of push rod 1340. The resulting bending moment about connection section 1330 can cause failure of adapter system 1300. To substantially reduce or eliminate asymmetrical loading, shoulder portion 1350 is preferably shaped to prevent such asymmetrical loading by, for example, being open on the top and bottom thereof (see, Figure 10B). Removing a bottom edge of abutment shoulder 1350 where cone region 240 of syringe 200 would otherwise rest results in generally symmetrical loading about the axis of adapter system 1300 (and syringe 200) and substantially reduces or removes lateral loads and bending moments

during forward plunger advancement. Axial load applied to the end of adapter assembly 1300 is maximized while lateral load is minimized.

In addition, enforcements such as a rib 1354 can be added to carrier section 1310 to limit wall flexing by increasing the material strength as illustrated in Figure 10B. Likewise, extensions 1556 and 1558 illustrated in Figure 12B can be formed integrally on the front portion of carrier section 1510 of adapter assembly 1500 (or adapter assembly 1300). Cone or transition region 240 of syringe 200 preferably does not contact extensions 1556 and 1558. When syringe 200 contacts shoulder portions 1350 (not shown in Figure 12B) with forward force, syringe 200 can act as a wedge and force the opposing portions of shoulder 1350 apart. Forming extensions 1556 and 1558 around the circumference of the front of carrier section 1510 creates hoop forces or stresses that help prevent such deformation. Extensions 1556 and 1558 are preferably spaced sufficiently at the top portion thereof to allow ready removal of syringe 200 and any attached tubing as discussed above.

In addition to a rotating retaining member 1360 (which operates generally in a manner as described for retaining member 460 of Figures 4A through 4C) or in lieu thereof, adapter system 1300 preferably includes a biasing retaining section 1370 illustrated in Figure 10C and 10D. In the embodiment of Figures 10C and 10D, retaining section 1370 includes two flexing retaining members 1372 and 1374 inset from the inside wall of carrier section 1310 to provide a pressure fit against syringe 200, thereby securing syringe 200 in place. Retaining section 1370 assists, for example, in retaining syringe 200 within carrier section 1310 when the injector head (not shown in Figures 10A-D) is rotated to a position other than horizontal when syringe 200 is placed with carriage section 1310. Should, for example, the injector head (and thereby carrier section 1310) be in a vertical orientation, retaining section 1370 prevents the syringe from falling out of carrier section 1310 even before retaining member 1360 can be rotated to a closed position.

In the embodiment of Figure 10A, retainer section 1310 is movably connected (for example, hingedly or even removably connected) to intermediate section 1320. By, for example, rotating carrier section 1310 downward so that syringe 200 can be pulled forward without contacting the forward portion of carrier 1310, syringe 200 can be released and removed without retracting the plunger thereof following a full or partial injection.

In the embodiment of Figure 10A, carrier section 1310 is hingedly attached to intermediate section 1320 via pin joints (not shown) on each side thereof. Carrier section 1310 rotates about such pin points when closing the adapter assembly until extending members 1370 and 1372 contact abutment surfaces 1374 on intermediate section 1320. At that point, intermediate section 1320 is generally aligned with carrier section 1310, and carrier section 1310 is in position for shoulder 1350 to abut syringe transition region 240. A locking ring 1380 is preferably slidably positioned on extending members 1370 and 1372. After carrier section 1310 is rotated to a closed position, locking ring 1380 is slid rearward to abut a flange 1390 formed on the lower half of the front of intermediate section 1310 to lock carrier section 1310 in a closed position.

Figure 11 illustrates another embodiment of an adapter system 1400 that is very similar to adapter system 1300. However, carrier section 1410 is not hingedly attached to intermediate section 1420. In this embodiment, push rod 1440 includes a movable section (for example, hinging or rotating section 1442) positioned forward of intermediate section 1420. Hinging section 1442 allows syringe 200 to be moved (rotated, in this embodiment) out of alignment with the axis of adapter system 1400 so that syringe 200 can be removed without retracting the drive member of the injector following a full or partial injection.

Figures 12A and 12B illustrate an adapter system 1500 that includes a carrier section 1510, an intermediate section 1520 and a connector section 1530 as discussed above. Adapter system 1500 includes a cleaning or contact member such as a wiper seal 1570 and retainer ring 1580 for positioning wiper seal 1570 within

adapter system 1500. Wiper seal 1570 operates to remove unwanted contrast media (resulting, for example, from leakage and/or spillage) from push rod 1540. In that regard, as the piston is being retracted following an injection, wiper seal 1570 cleans/wipes any contrast media that has inadvertently adhered to push rod 1540  
5 therefrom. Moreover, wiper seal 1570 also minimizes unwanted contrast media from entering the intermediate section 1520 of the adapter system 1520. Retainer 1580 preferably holds wiper seal 1570 in place via a pressure fit in carrier section 1510.

As best illustrated in Figure 12B, push rod 1540 is preferably prevented from being removed from the rear opening of connection section 1330 by a  
10 retainer flange 1590 that abuts a flange on the rearward end of push rod 1540 (not shown in Figure 12B; see push rod 1640 in Figure 13B). The flange on the rearward end of push rod 1540 is preferably dimensioned to be slightly smaller than the inner diameter of intermediate section 1520. The contact of contact member 1570 with push rod 1540 and the contact of the push rod flange with the inner wall of  
15 intermediate section 1520 assist in maintaining proper alignment of push rod 1540 within adapter assembly 1500.

Figures 13A and 13B illustrate an adapter system 1600 including a carrier section 1610, an intermediate section 1620, a connector section 1630 and a pushrod 1640. Adapter system 1600 includes a forward-biasing, rear abutment  
20 member 1670 that operates to bias syringe 200 completely forward in carrier section 1610 regardless of the length of syringe 200. Abutment member 1670 is preferably slidably positioned at the rear of carrier section 1610 and is biased forward by, for example, a spring 1674 housed within intermediate section 1620. As the operator loads a syringe into carrier section 1610, by initially placing bottom  
25 flange 230 of syringe 200 against abutment member 1670, spring 1674 is compressed as required to accommodate various syringe lengths. With spring 1674 compressed, syringe 200 is biased completely forward within carrier section 1610. Abutment member 1670 also preferably includes a generally central passage 1676 formed therethrough to allow pushrod 1640 to impart force to the syringe plunger.

Figure 14 illustrates another forward biasing abutment member 1770 suitable for use in an adapter system 1700 to bias syringe 200 completely forward within carrier section 1710. Abutment member 1770 is slidably seated within carrier section 1710 and includes flex members 1772 that abut, for example, a forward facing surface 1712 of carrier section 1710 and bias abutment member 1770 forward. Abutment member 1770 of Figure 14 includes three sets of inward projecting protrusions 1774 for contacting rear flange 230 of syringe 200. Multiple sets of protrusions 1774 are provided to accommodate multiple syringe lengths. Flex members 1772 push against surface 1712 when syringe 200 is in place, thereby biasing syringe 200 in a fully forward position within carrier section 1710 regardless of syringe length.

Although the present invention has been described in detail in connection with the above examples, it is to be understood that such detail is solely for that purpose and that variations can be made by those skilled in the art without departing from the spirit of the invention except as it may be limited by the following claims.

## WHAT IS CLAIMED IS:

1. An adapter for releasably attaching a syringe to a front-loading powered injector, wherein the syringe comprises a body, a transition region attached to a forward end of the body and a plunger slideably positioned within the body, and wherein the injector comprises a front wall, an opening formed in the front wall, and a drive member reciprocally mounted in the injector, the adapter comprising:
  - a syringe carrier adapted to seat at least a portion of the syringe, the syringe carrier comprising at least one rearward facing abutment member to abut at least one forward facing surface of the transition region, the syringe carrier comprising an opening therein to allow the drive member of the injector to communicate forward force to the plunger through abutment without connective engagement between the drive member and the plunger; and
  - a releasable mounting mechanism positioned to the rear of the syringe carrier to mount the adapter in a desired position relative to the front wall of the injector.
3. The adapter of Claim 1 wherein the abutment member abuts the transition region in the vicinity of the transition from the body to the transition region.
6. The adapter of Claim 1 further comprising a first section and a second section, the first section and the second section being rotatable relative to each other about a hinge axis generally perpendicular to a longitudinal axis of the adapter, the first section and the second section being rotatable about the hinge axis to an open position to allow loading of the syringe into the adapter from a position to the rear of the hinge axis, the first section and the second section also being rotatable about the hinge axis to a closed position to form the syringe carrier.
7. The adapter of Claim 1 further comprising a first section and a second section, the first section and the second section being generally the same in construction, the first section and the second section being connectable to form the syringe carrier and the releasable mounting mechanism.

8. An adapter for releasably attaching a syringe to a powered injector including a drive member reciprocally mounted therein, the adapter comprising:

a first section and a second section, the first section and the second section being rotatable relative to each other about a hinge axis generally perpendicular to a longitudinal axis of the adapter, the first section and the second section being rotatable about the hinge axis to an open position to allow loading of the syringe into the adapter from a position to the rear of the hinge axis, the first section and the second section also being rotatable about the hinge axis to a closed position to form a syringe carrier to seat at least a portion of the syringe.

9. The adapter of Claim 8 wherein the powered injector further includes a front wall, and the adapter further includes a releasable mounting mechanism positioned to the rear of the syringe carrier to mount the adapter in a desired position relative to the front wall of the injector.

10. An adapter for releasably attaching a syringe to a powered injector including a drive member reciprocally mounted therein, the adapter comprising:

a first section and a second section, the first section and the second section being generally the same in construction, the first section and the second section being connectable to form a syringe carrier to seat at least a portion of the syringe.

11. The adapter of Claim 10 wherein the injector further includes a front wall, and the first section and the second section also form a releasable mounting mechanism positioned to the rear of the syringe carrier to mount the adapter in a desired position relative to the front wall of the injector.

12. The adapter of Claim 6, further comprising a releasable latch to maintain the first section and the second section in the closed position.

13. The adapter of Claim 1 wherein at least a portion of the adapter is formed to enable viewing of the syringe.

14. The adapter of Claim 1 wherein the releasable mounting mechanism comprises flanges that cooperate with the injector to mount the adapter to the  
5 injector.

15. The adapter of Claim 1, further comprising a sealing flange positioned forward of the mounting mechanism to abut the front wall of the injector.

16. The adapter of Claim 1, further comprising indicia to provide information to the injector.

10 17. The adapter of Claim 1, further comprising at least one syringe abutment member to prevent the syringe from rotating within the adapter.

18. The adapter of Claim 6 wherein the hinge axis is off center from a center line of the adapter so that force experienced during an injection tends to maintain the first section and the second section in the closed position.

15 19. The adapter of Claim 1 wherein the syringe further comprises a handle attached to the plunger, the handle extending out of the rear of the syringe body, the adapter further comprising a guide member to guide the movement of the handle of the syringe during an injection.

20 20. The adapter of Claim 3 wherein the abutment member comprises a resilient material to abut the transition region in the vicinity of the transition from the body to the transition region.

21. The adapter of Claim 1 wherein the carrier is formed to have an open state such that the top thereof is open along the length of the carrier.



22. The adapter of Claim 21, further comprising at least one syringe retaining mechanism for retaining the syringe within the carrier during an injection.

23. The adapter of Claim 22 wherein the retaining mechanism is rotatable around an axis of the adapter between an open position and a closed position.

5           24. The adapter of Claim 22 wherein the retaining mechanism is slidable along an axis of the adapter.

25. The adapter of Claim 22 wherein the retaining mechanism comprises at least one extending member that is biased in a position to retain the syringe in the adapter.

10           26. The adapter of Claim 22 wherein the retaining mechanism is hingedly attached to one side of the adapter to be movable between an open position and a closed position.

15           27. The adapter of Claim 1 wherein the releasable mounting mechanism comprises a plate that is removably attachable to the front wall of the injector.

28. The adapter of Claim 9 wherein the releasable mounting mechanism comprises flanges that cooperate with the injector to mount the adapter on the injector.

20           29. The adapter of Claim 9, further comprising a sealing flange positioned forward of the mounting mechanism to abut the front wall of the injector.

30. The adapter of Claim 8, further comprising indicia to provide information to the injector.

31. The adapter of Claim 8, further comprising at least one syringe abutment member to prevent the syringe from rotating within the adapter.

32. The adapter of Claim 8 wherein the hinge axis is off center from a center line of the adapter so that force experienced during an injection tends to maintain the first section and the second section in the closed position.

33. The adapter of Claim 8 wherein the syringe further comprises a plunger disposed within the syringe body and a handle attached to the plunger extending out of the rear of the syringe body, the adapter further comprising a guide member to guide the movement of the handle of the syringe during an injection.

34. The adapter of Claim 8, further comprising a releasable latch to maintain the first section and the second section in the closed position.

35. The adapter of Claim 8 wherein the syringe further comprises a syringe flange attached to a rearward end of the body, the adapter further comprising an abutment member to abut a forward facing surface of the syringe flange.

36. The adapter of Claim 35 wherein the abutment member abuts the syringe flange in the vicinity of the transition from the body to the syringe flange.

37. The adapter of Claim 36 wherein the abutment member comprises a resilient material to abut the syringe flange in the vicinity of the transition from the body to the syringe flange.

38. The adapter of Claim 8 wherein the syringe further comprises a transition region attached to a forward end of the body, the adapter further comprising an abutment member to abut a forward facing surface of the transition region.

39. The adapter of Claim 38 wherein the abutment member abuts the transition region in the vicinity of the transition from the body to the transition region.

40. The adapter of Claim 39 wherein the abutment member comprises a resilient material to abut the syringe flange in the vicinity of the transition from the  
5 body to the transition region.

41. The adapter of Claim 8 wherein at least a portion of the adapter is formed to enable viewing of the syringe.

42. The adapter of Claim 9 wherein the releasable mounting mechanism comprises a plate that is removably attachable to the front wall of the  
10 injector.

43. The adapter of Claim 11 wherein the releasable mounting mechanism comprises flanges that cooperate with the injector to mount the adapter on the injector.

44. The adapter of Claim 43, further comprising a sealing flange  
15 positioned forward of the mounting mechanism to abut the front wall of the injector.

45. The adapter of Claim 43 wherein the cooperation between the releasable mounting mechanism and the injector maintains the first section and the second section in a closed relationship.

46. The adapter of Claim 10, further comprising indicia to provide  
20 information to the injector.

47. The adapter of Claim 10, further comprising at least one syringe abutment member to prevent the syringe from rotating within the adapter.

48. The adapter of Claim 10 wherein at least a portion of the adapter is formed to enable viewing of the syringe.

49. The adapter of Claim 10 wherein the first section and the second section are hingedly attached along the sides thereof to be movable between an open position and a closed position.

50. The adapter of Claim 10 wherein the first section and the second section are rotatable relative to each other about a hinge axis generally perpendicular to a longitudinal axis of the adapter.

51. The adapter of Claim 50 wherein the syringe further comprises a syringe flange attached to a rearward end of the body, the adapter further comprising an abutment member to abut a forward facing surface of the syringe flange.

52. The adapter of Claim 51 wherein the abutment member abuts the syringe flange in the vicinity of a transition from the body to the syringe flange.

53. The adapter of Claim 52 wherein the abutment member comprises a resilient material to abut the syringe flange in the vicinity of the transition from the body to the syringe flange.

54. The adapter of Claim 50 wherein the syringe further comprises a transition region attached to a forward end of the body, the adapter further comprising an abutment member to abut a forward facing surface of the transition region.

55. The adapter of Claim 54 wherein the abutment member abuts the transition region in the vicinity of the transition from the body to the transition region.

56. The adapter of Claim 55 wherein the abutment member comprises a resilient material to abut the transition in the vicinity of the transition from the body to the transition region.

57. The adapter of Claim 11 wherein the releasable mounting  
5 mechanism comprises a plate that is removably attachable to the front wall of the injector.

58. An adapter for releasably attaching a syringe to a front-loading  
powered injector, wherein the syringe comprises a body, a syringe flange attached to a  
rearward end of the body, and wherein the injector comprises a front wall, an opening  
10 formed in the front wall, and a drive member reciprocally mounted in the injector, the  
adapter comprising:

a syringe carrier adapted to seat at least a portion of the syringe, the syringe carrier  
including at least one rearward facing abutment member operable to abut the syringe flange  
only in the vicinity of the transition from the body to the syringe flange, the syringe carrier  
15 comprising an opening therein to allow the drive member of the injector to communicate  
with an interior of the syringe to pressurize fluid within the syringe; and

a releasable mounting mechanism positioned to the rear of the syringe carrier to  
mount the adapter in a desired position relative to the front wall of the injector.

59. The adapter of Claim 58 wherein the abutment member comprises a  
20 resilient material to abut the syringe flange in the vicinity of the transition from the body to  
the syringe flange.

1/43

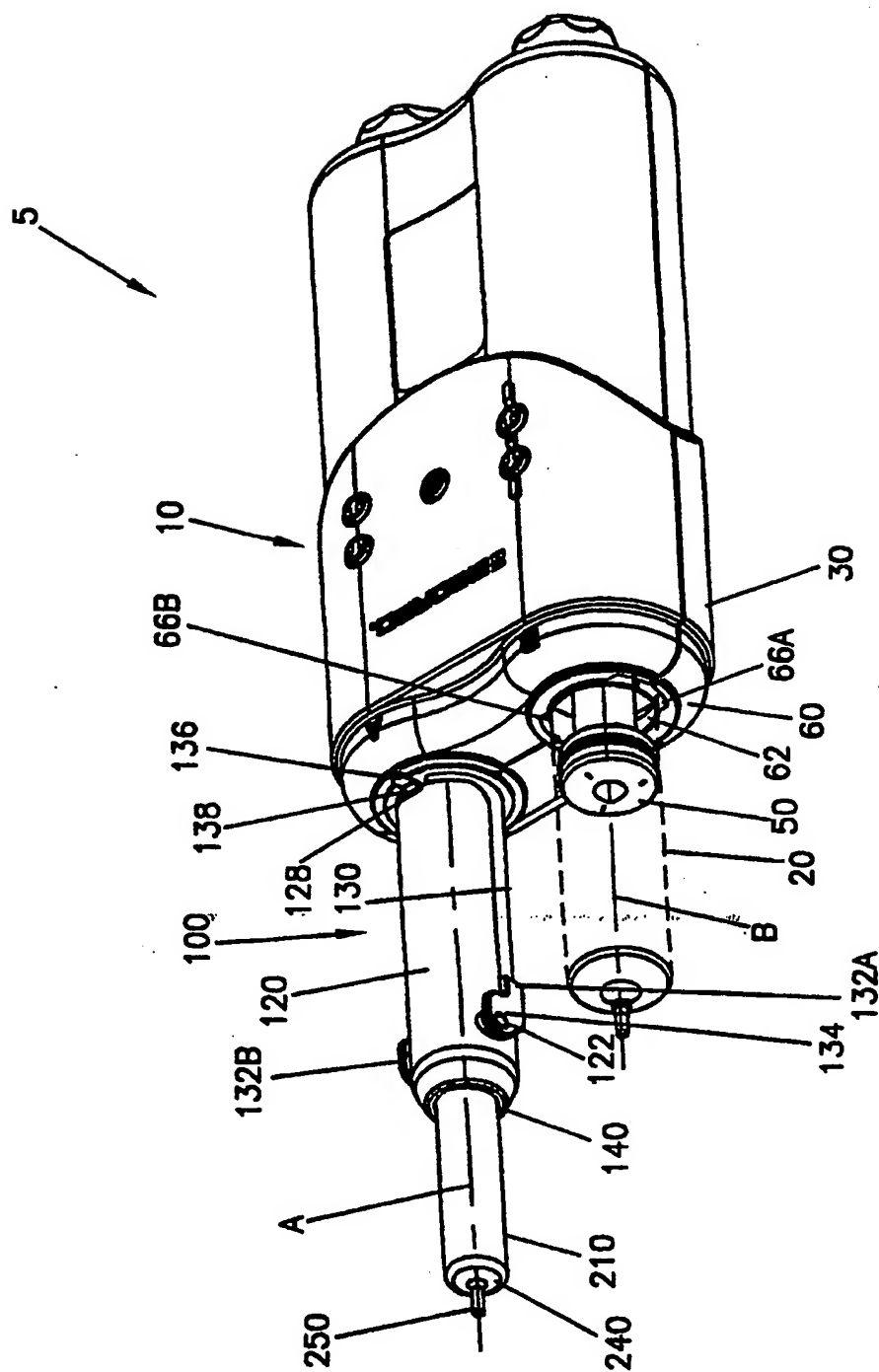


FIG. 1A



3/43

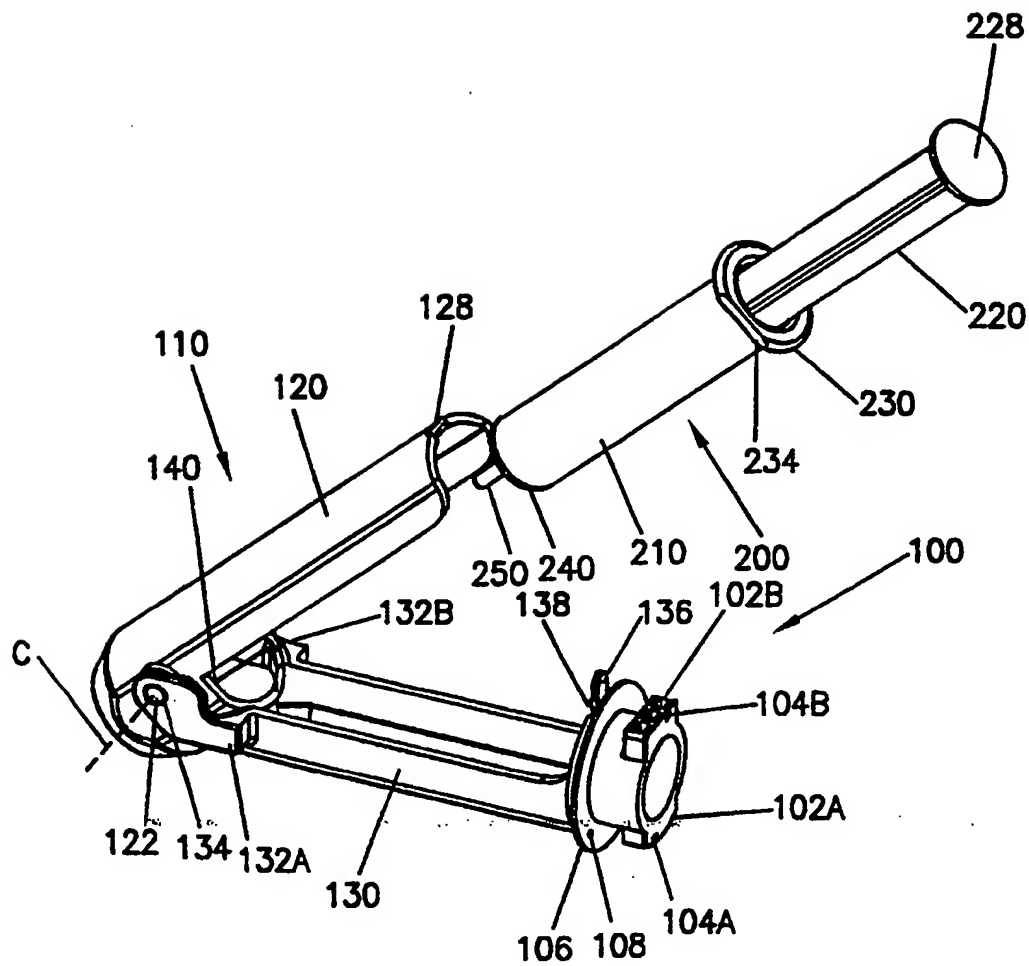


FIG. 2A



4/43

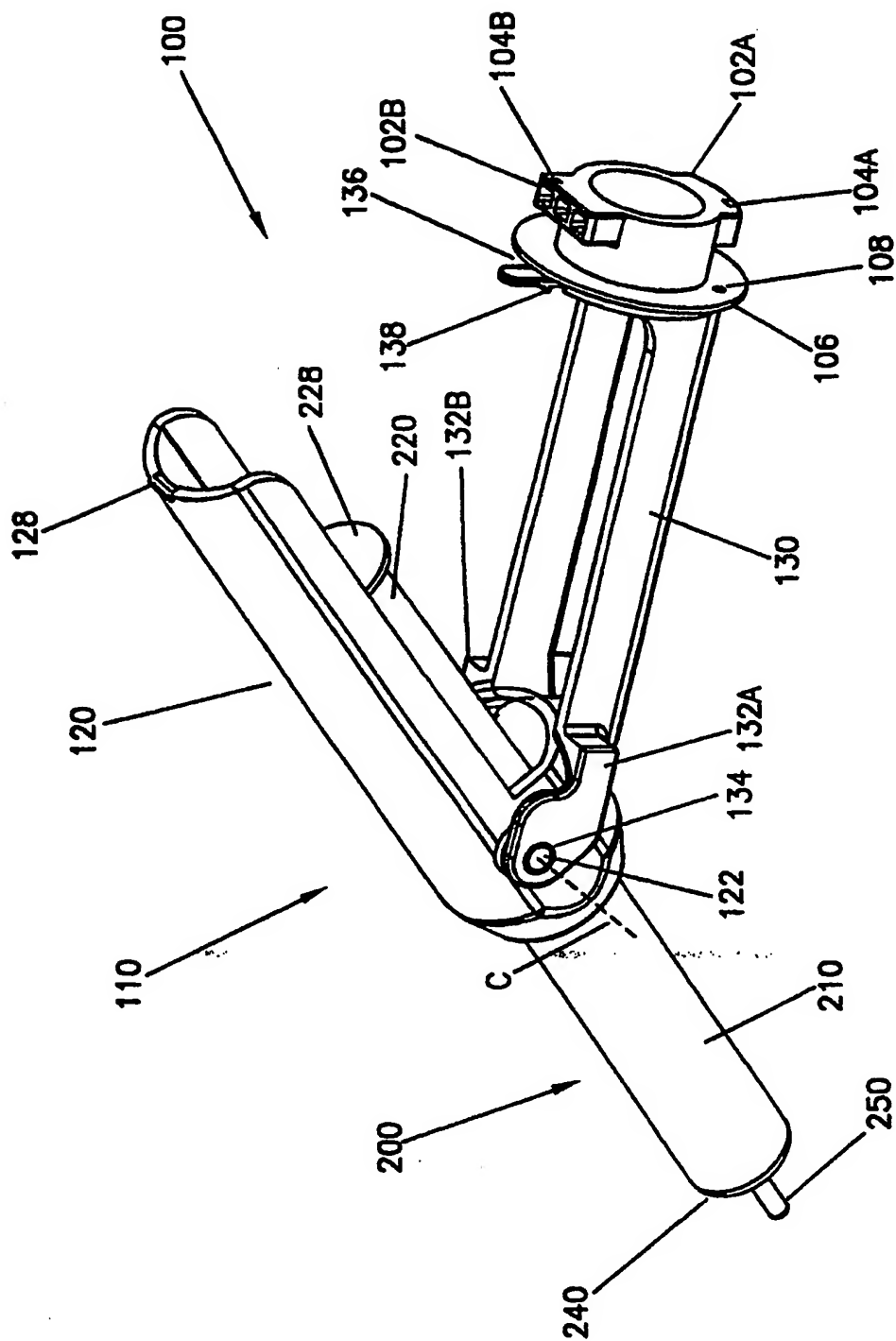


FIG. 2B

5/43

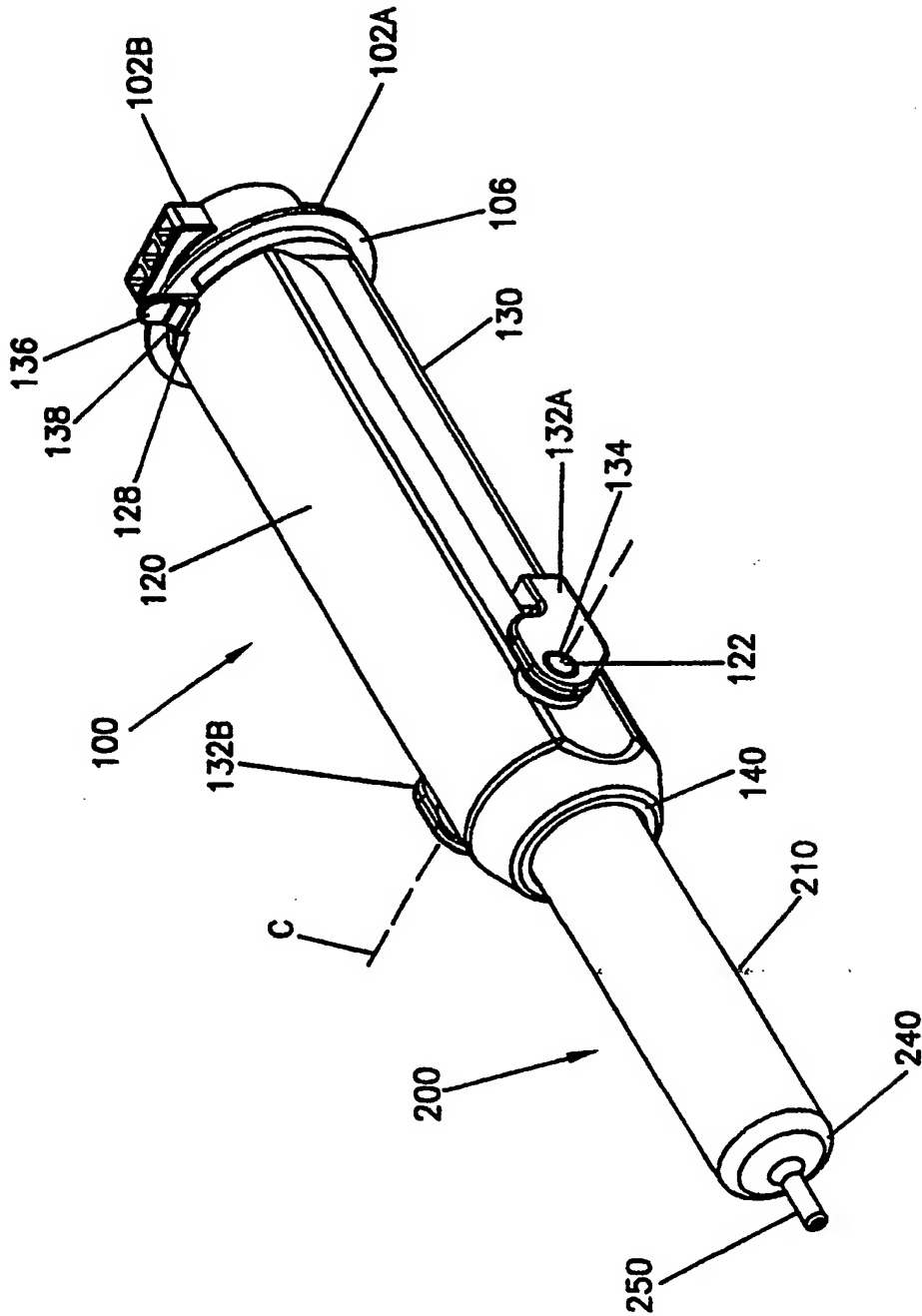


FIG. 2C

6/43

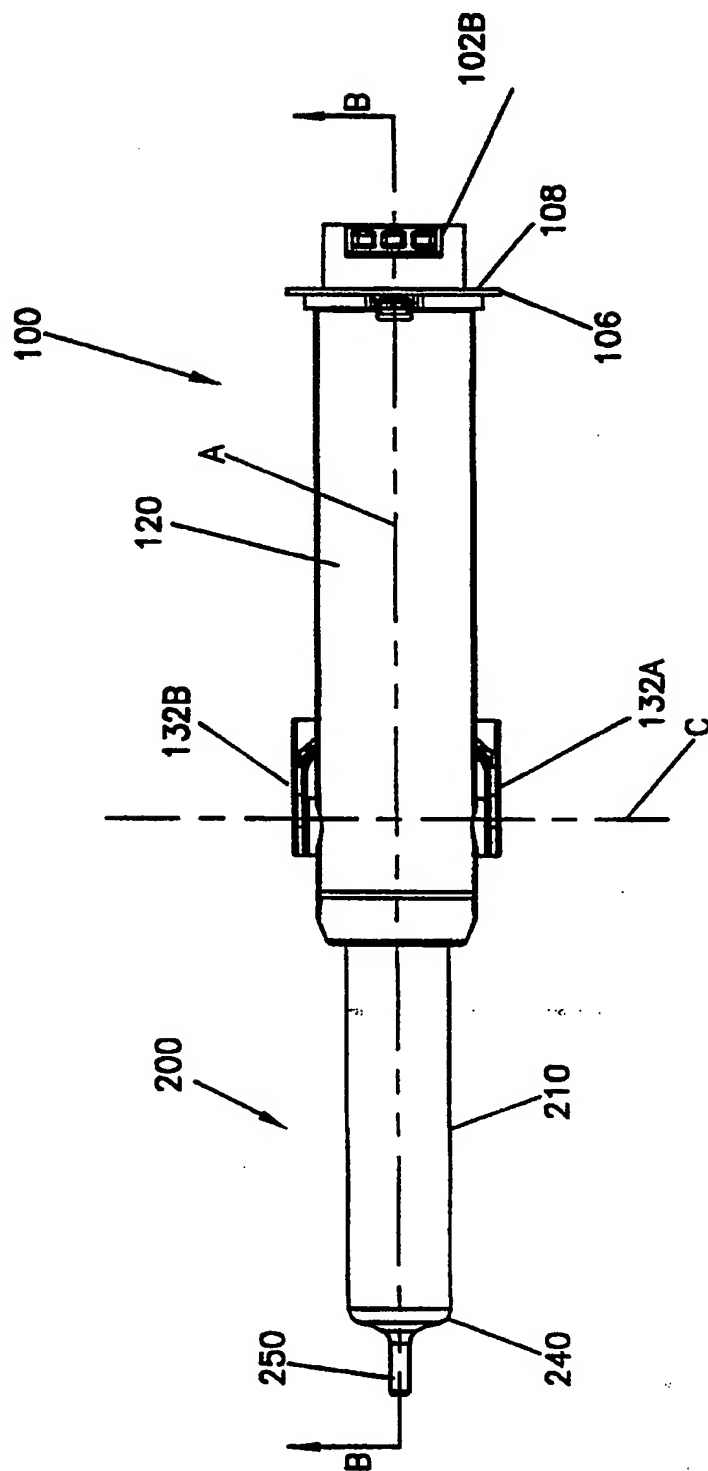


FIG. 2D

7/43

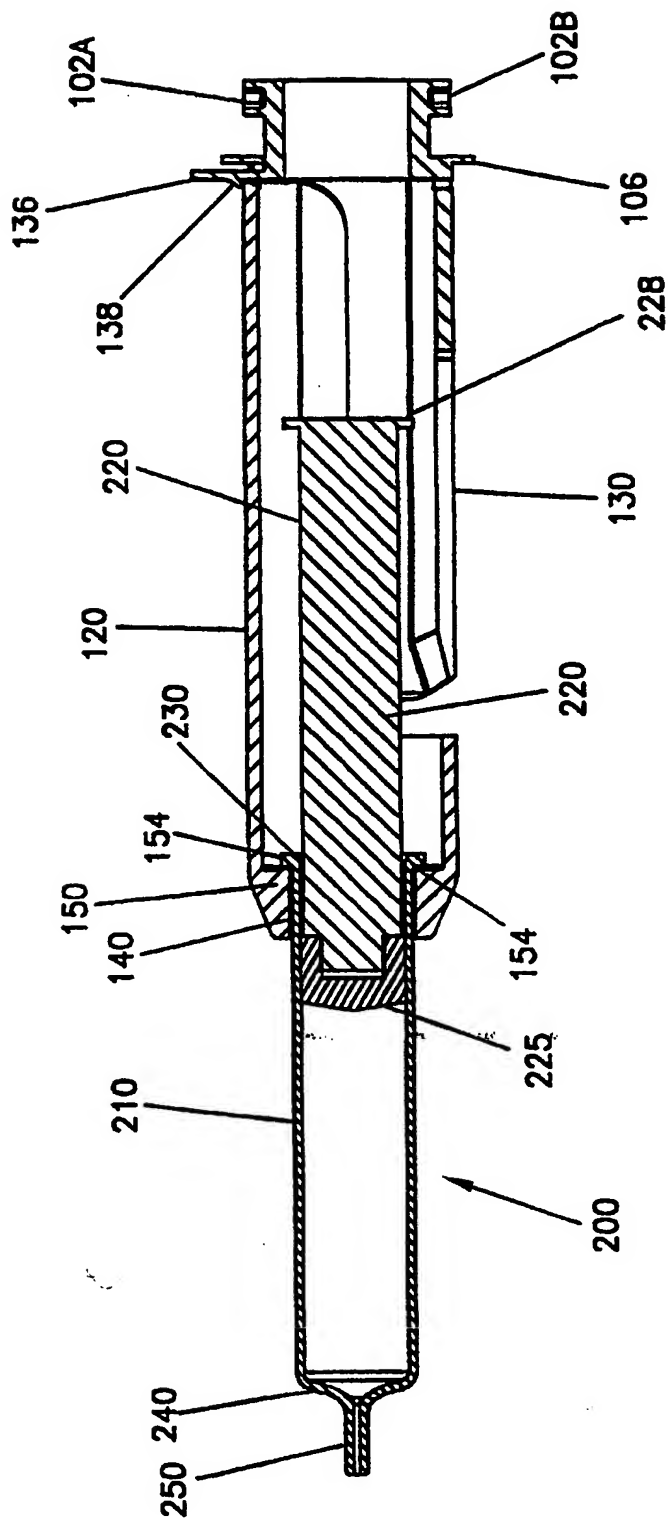


FIG. 2E



9/43

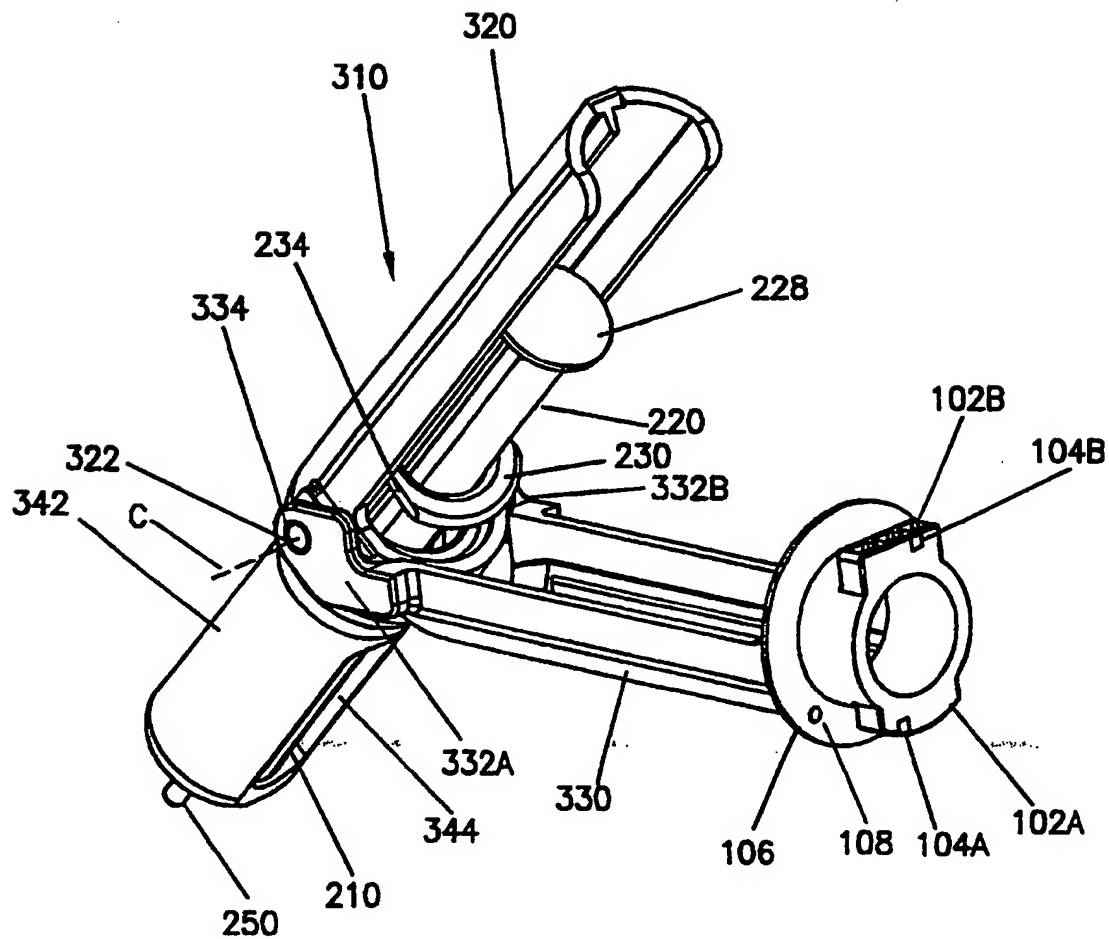


FIG. 3B

10/43

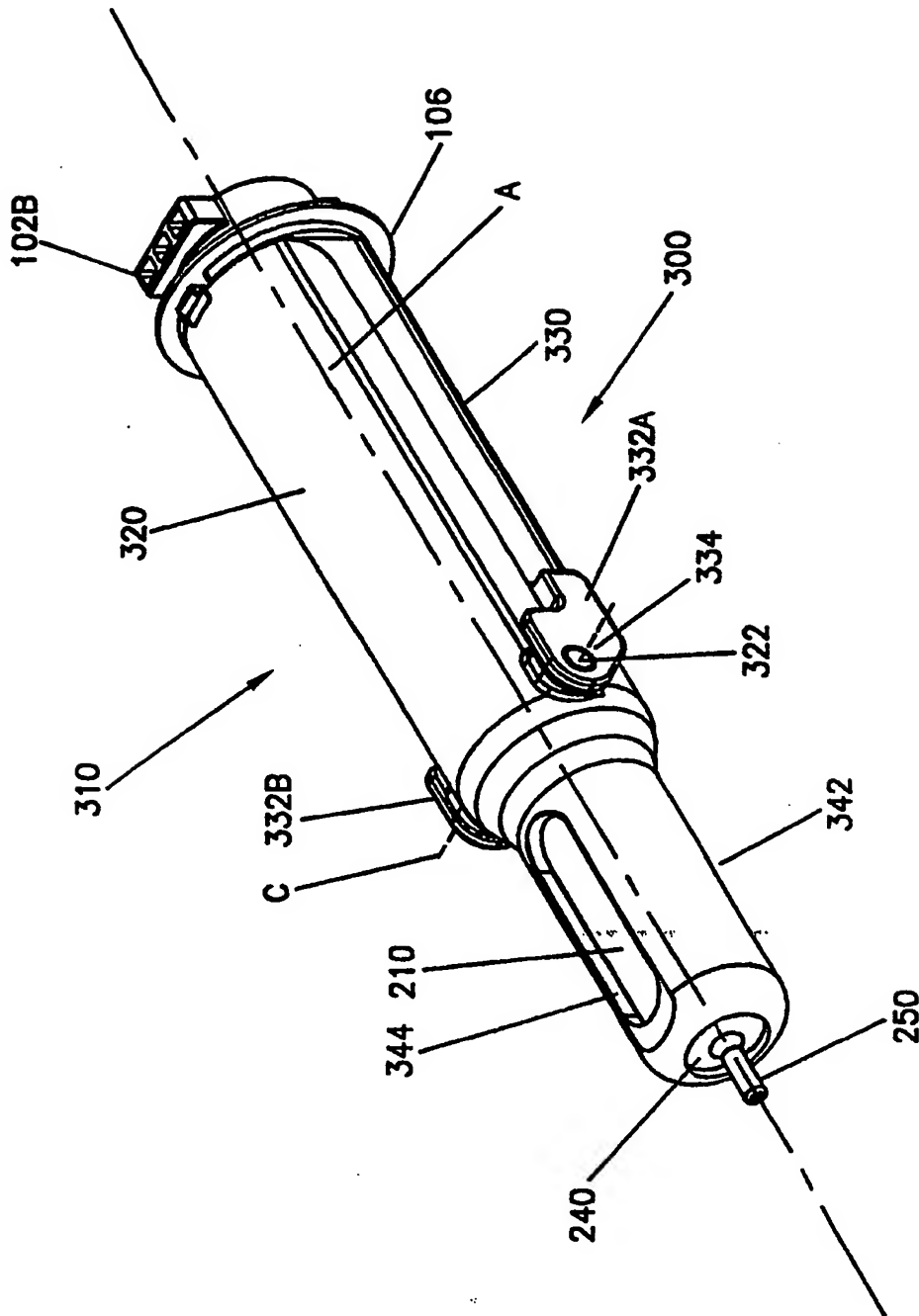


FIG. 3C

11/43

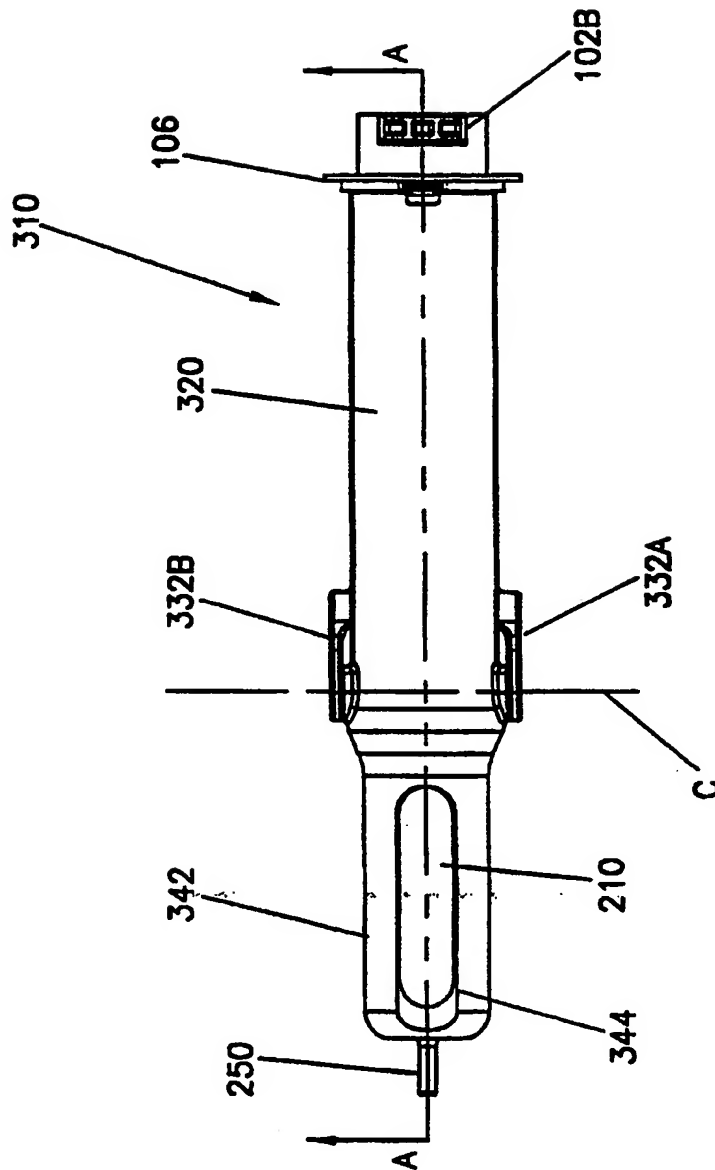


FIG. 3D







14/43

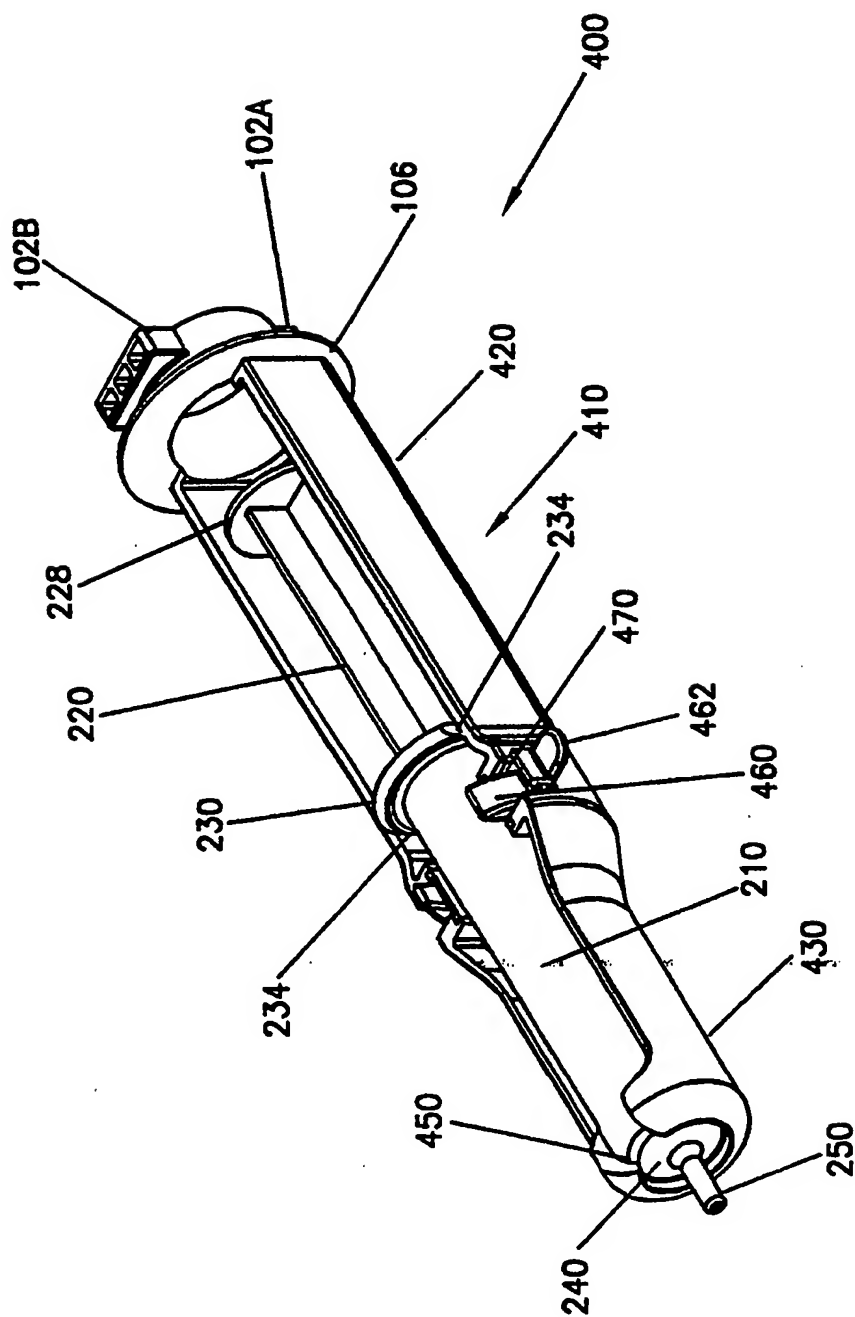


FIG. 4B



16/43

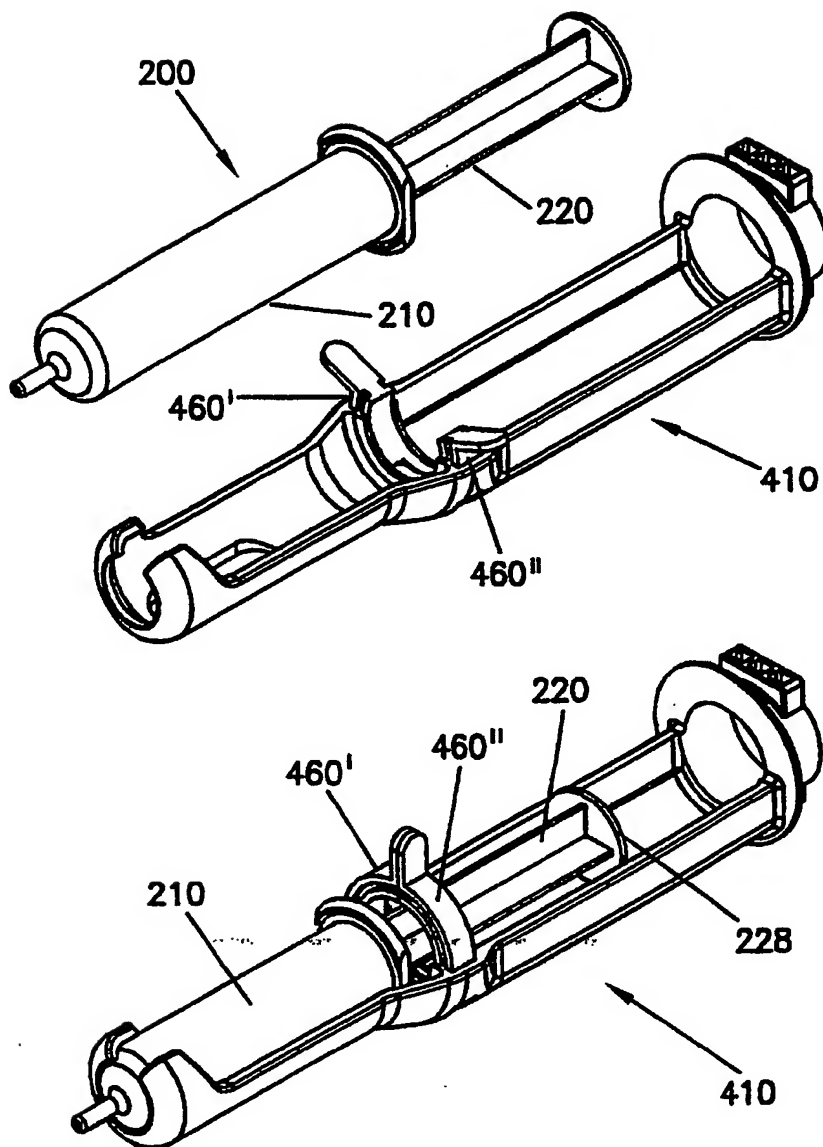


FIG. 4D

17/43

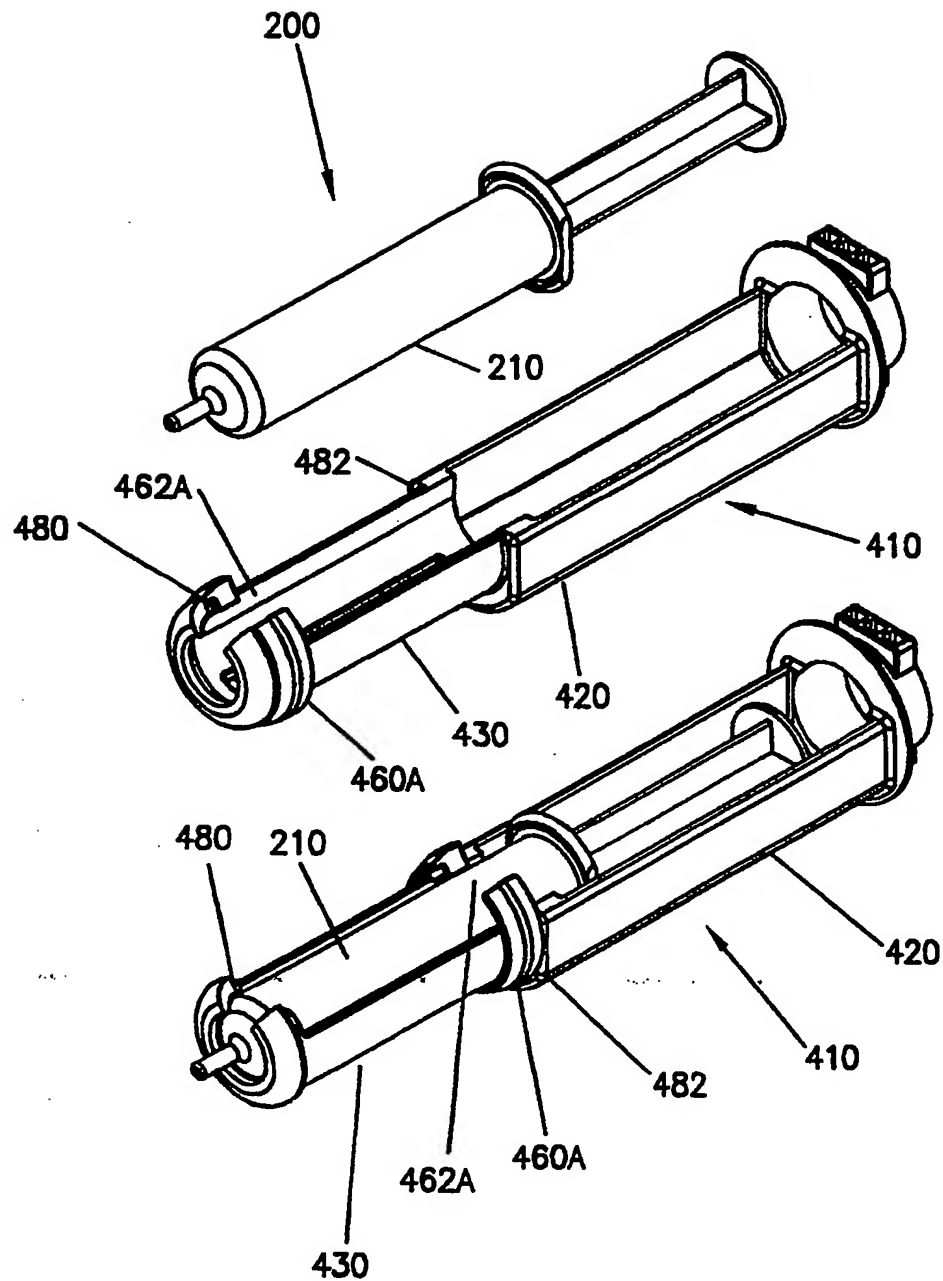


FIG. 4E

18/43

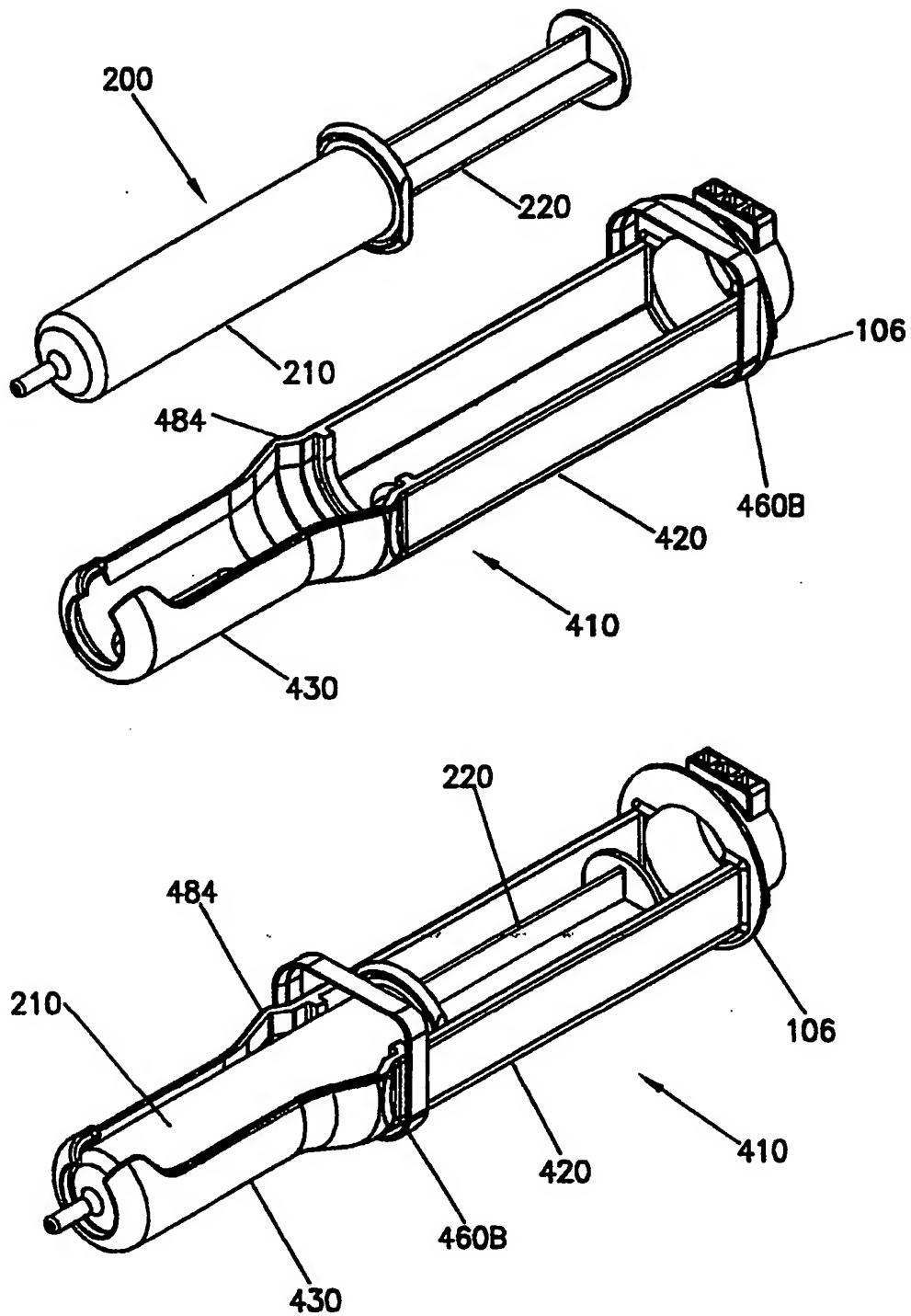


FIG. 4F

19/43

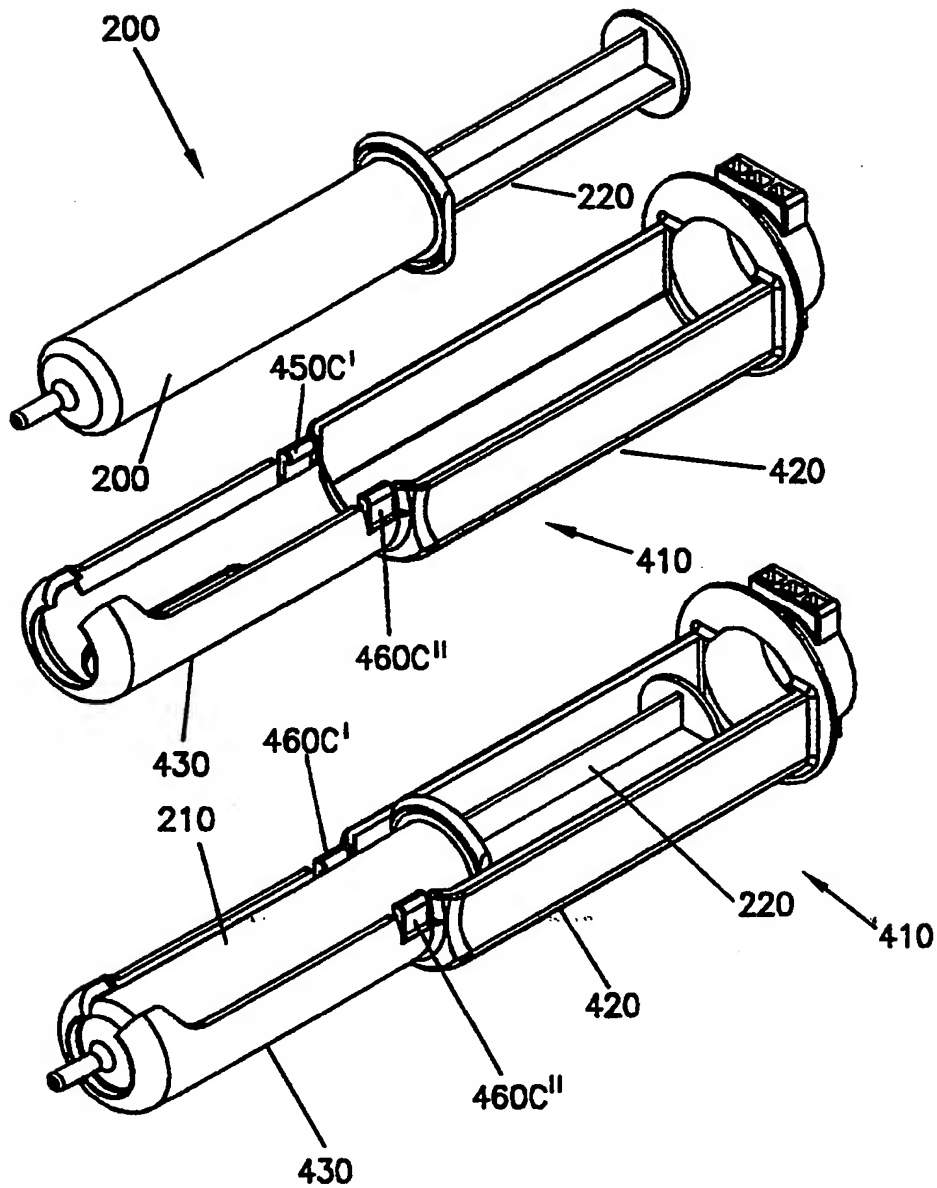


FIG. 4G



20/43

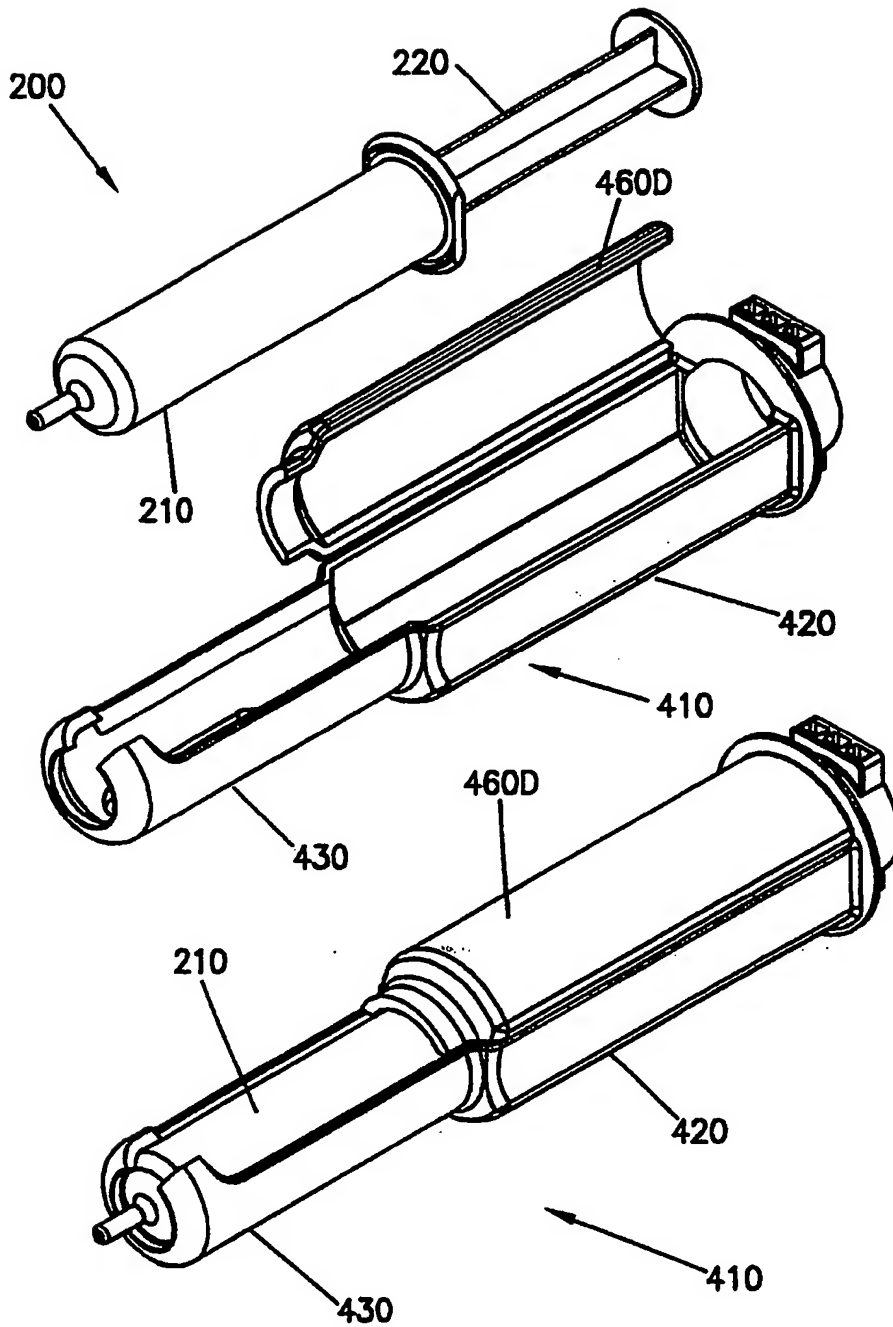


FIG. 4H



22/43

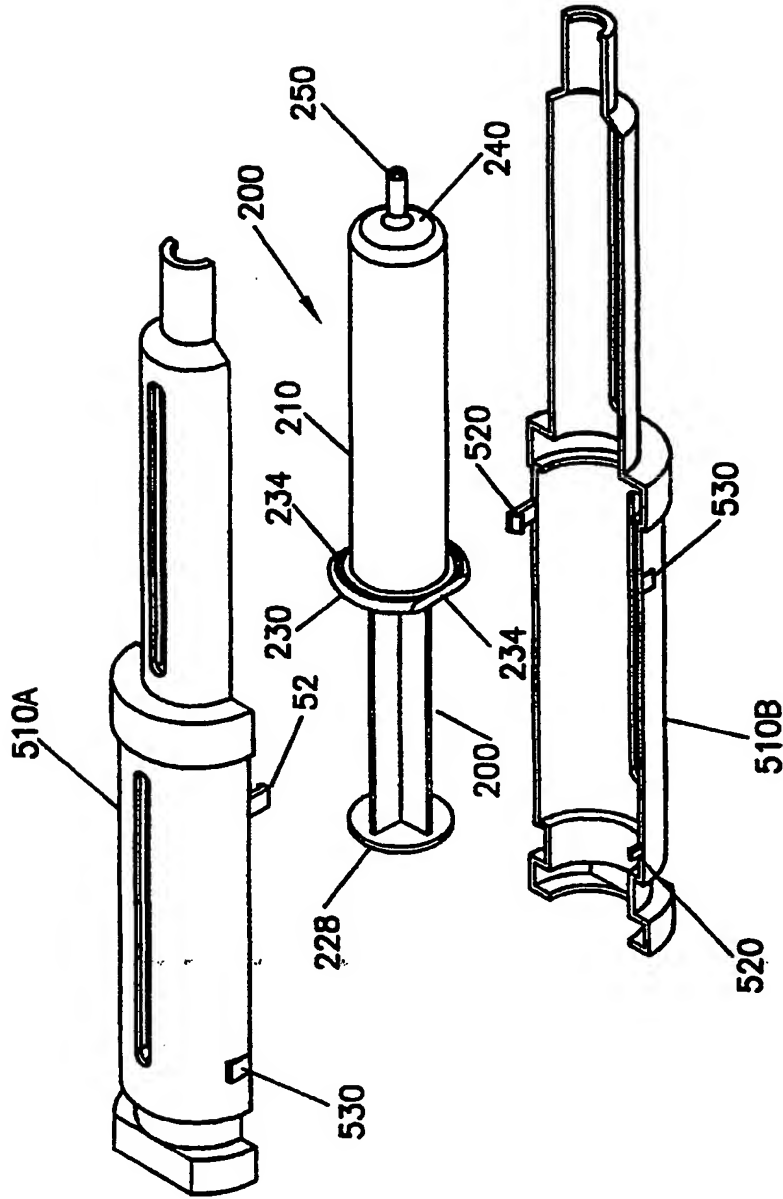


FIG. 5A

23/43

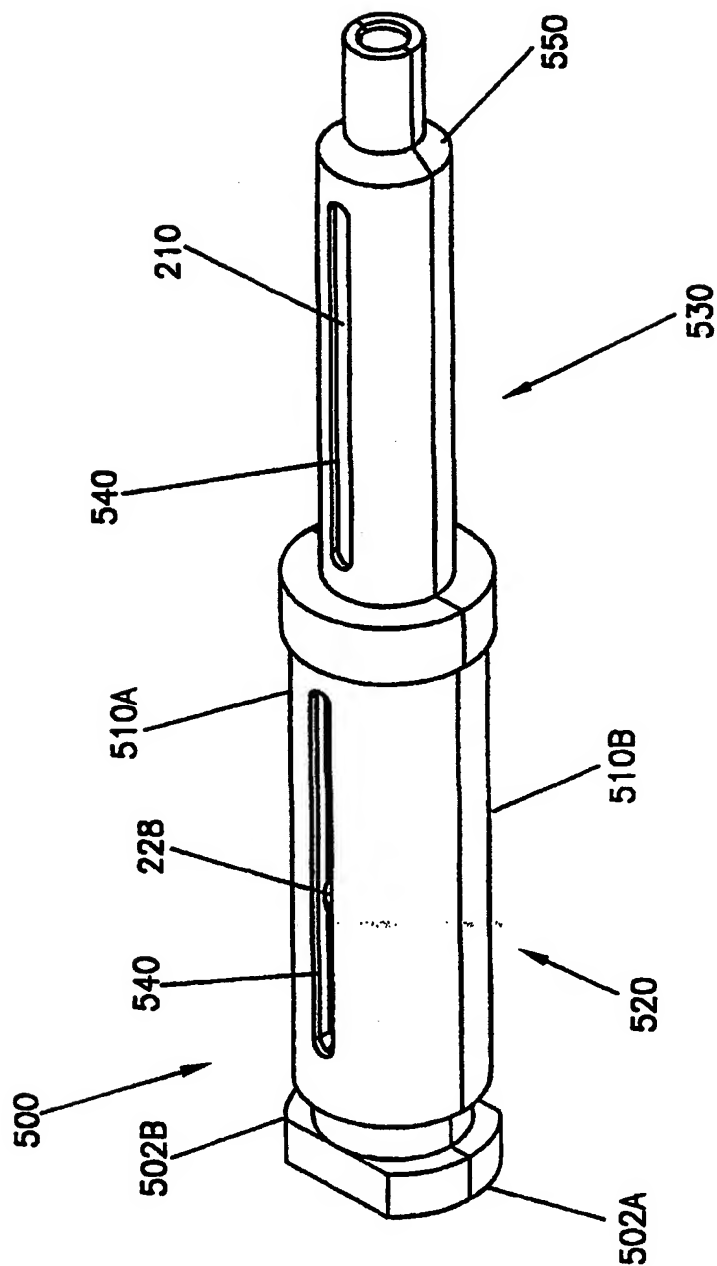


FIG. 5B

24/43

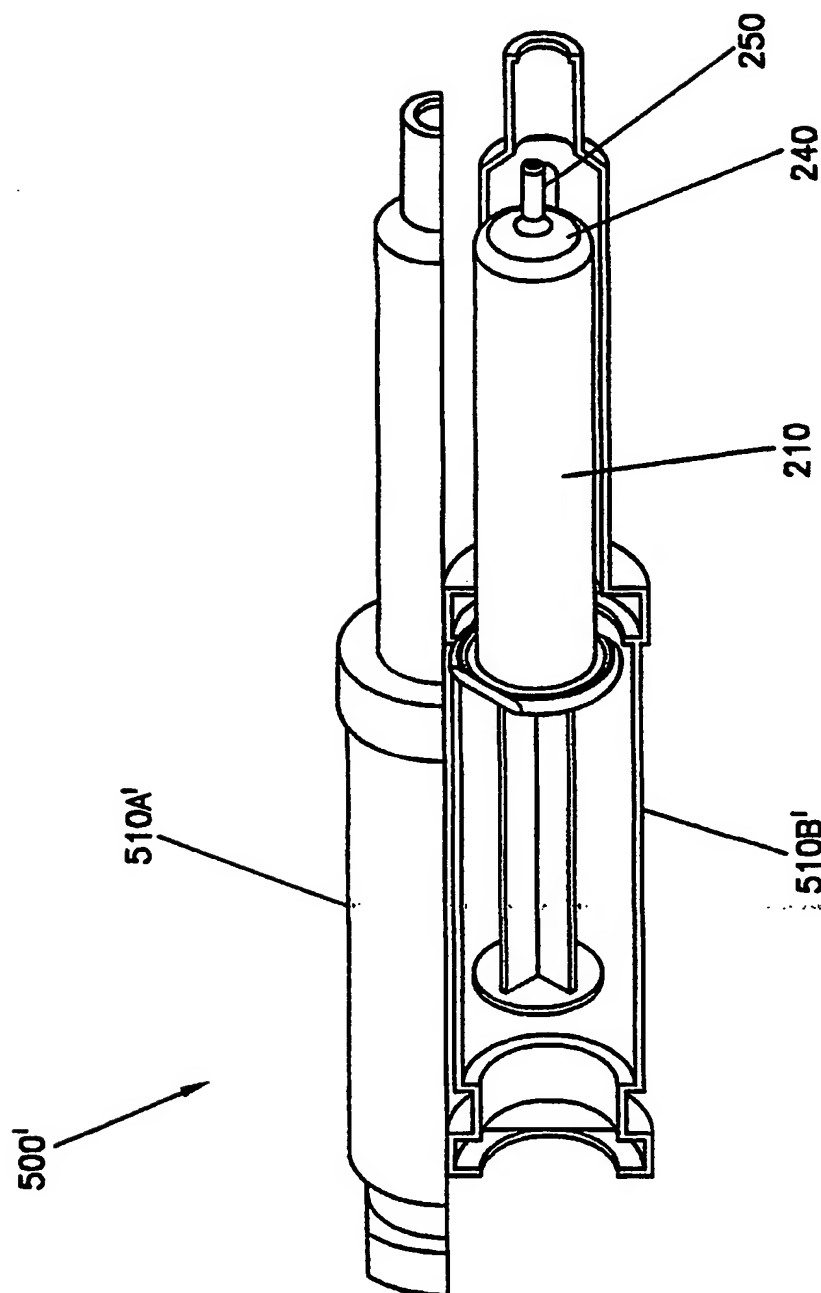


FIG. 6A

25/43

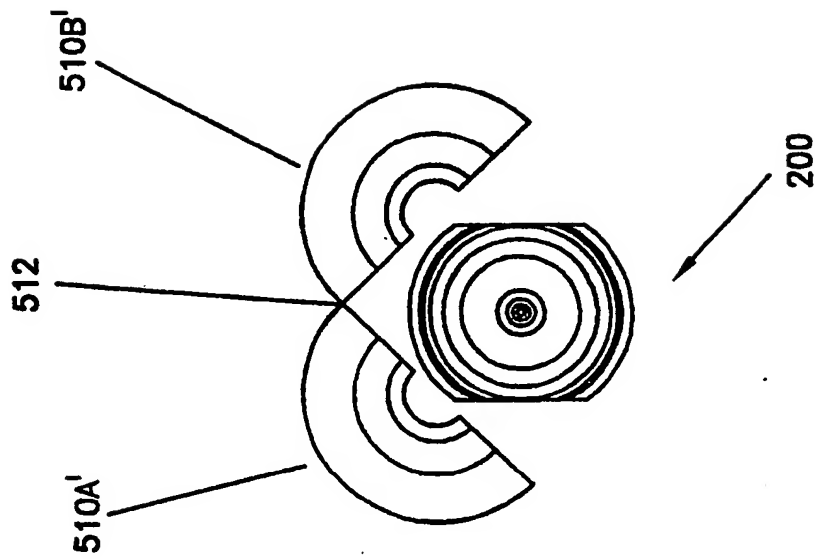


FIG. 6B

26/43

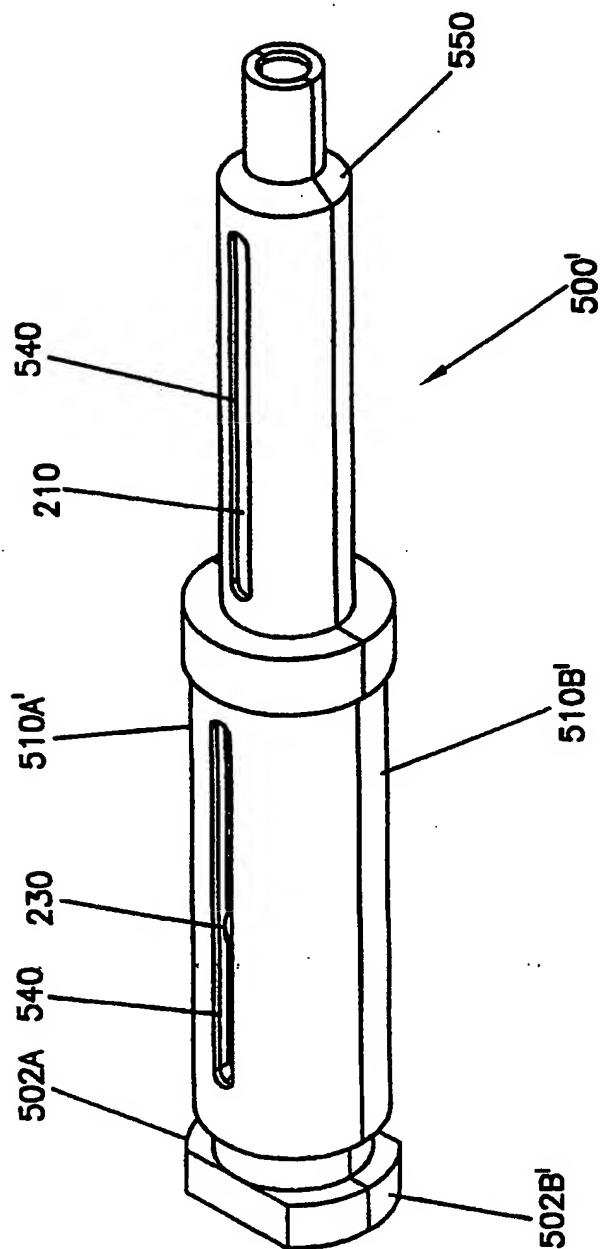


FIG. 6C

27/43

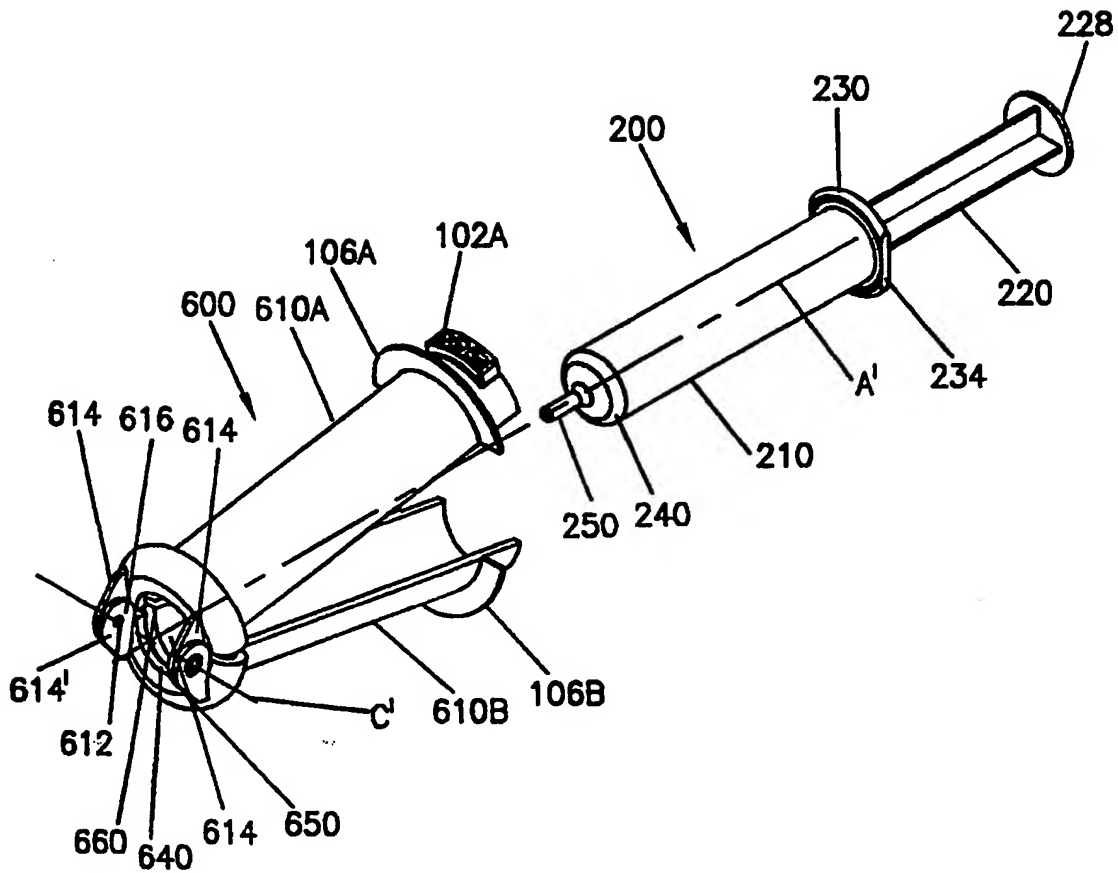


FIG. 7A



28/43

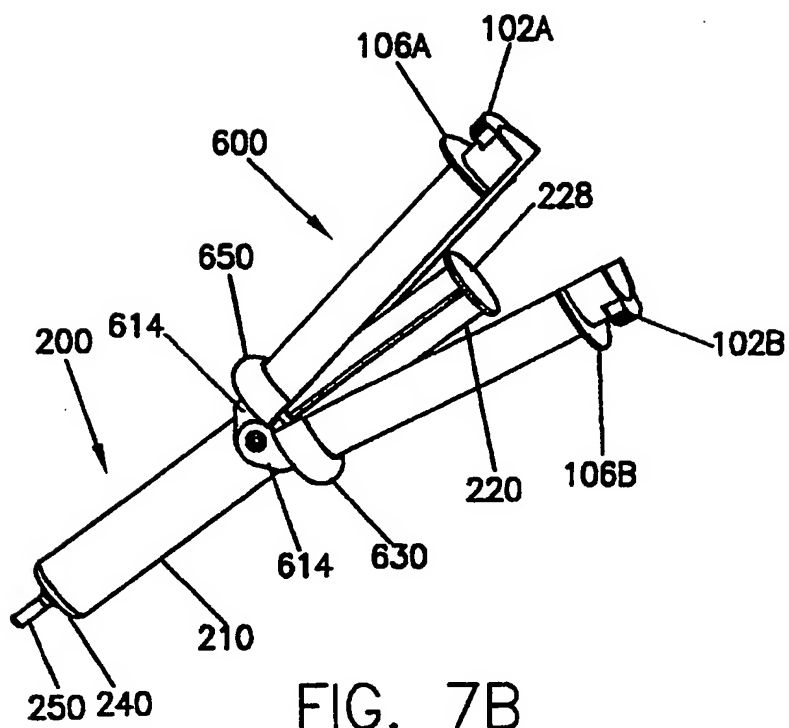


FIG. 7B

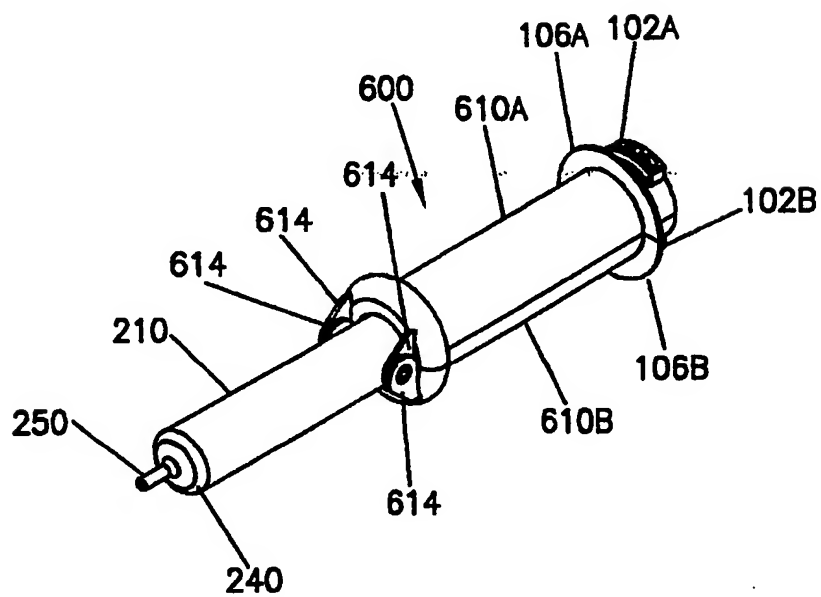


FIG. 7C

29/43

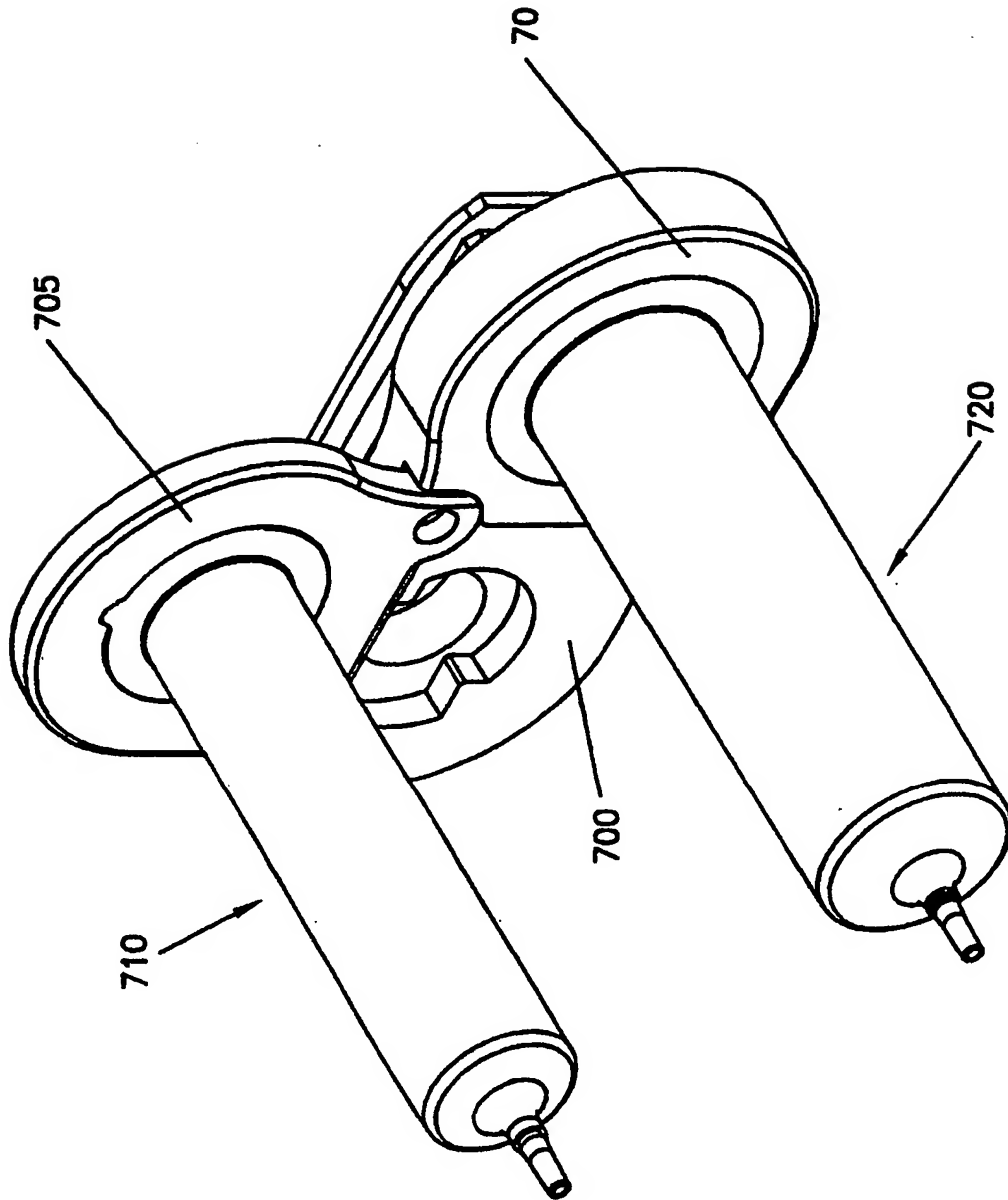


FIG. 8A

30/43

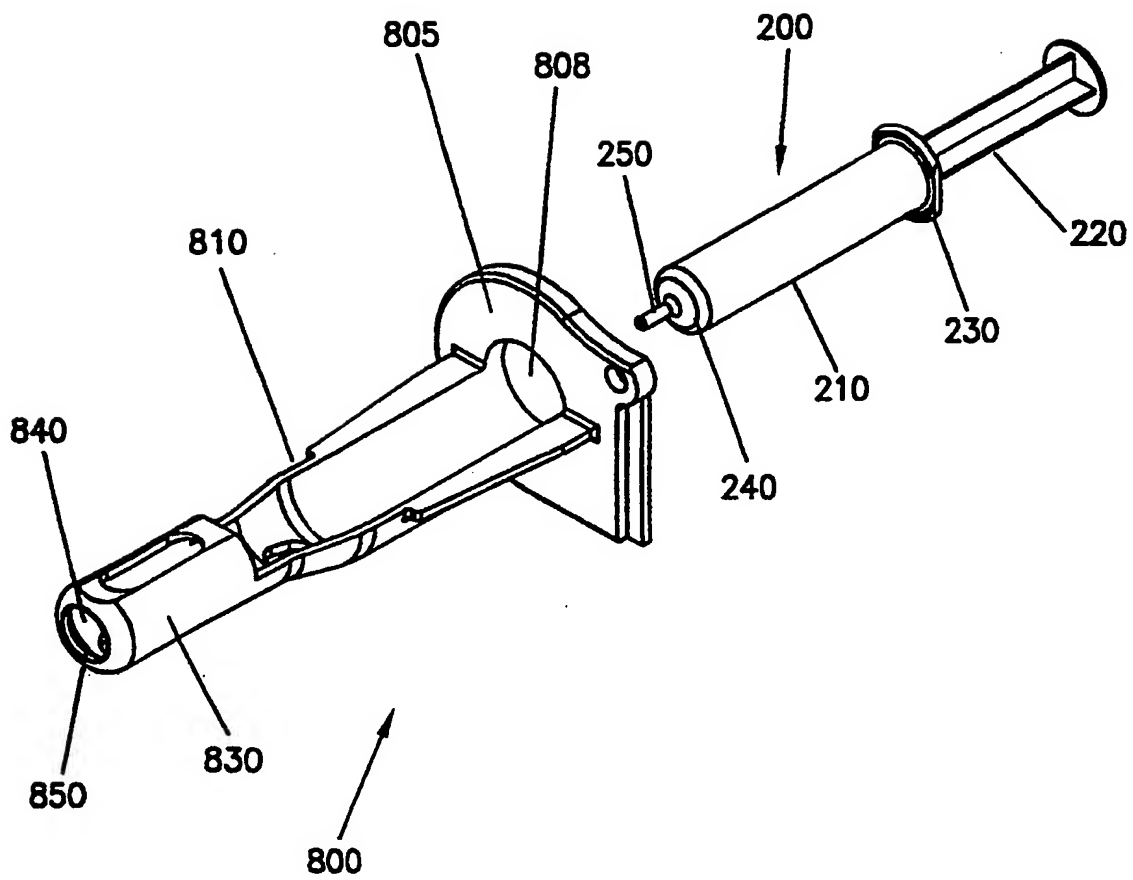


FIG. 8B

31/43

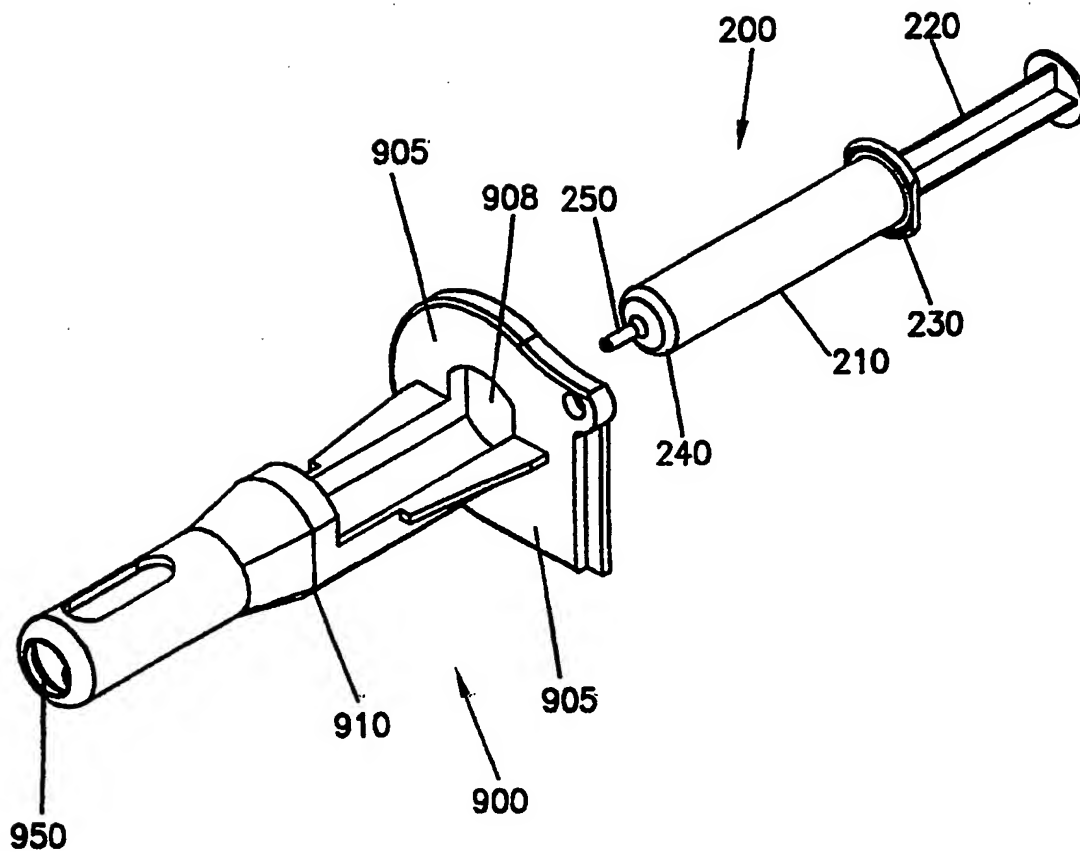


FIG. 8C

32/43

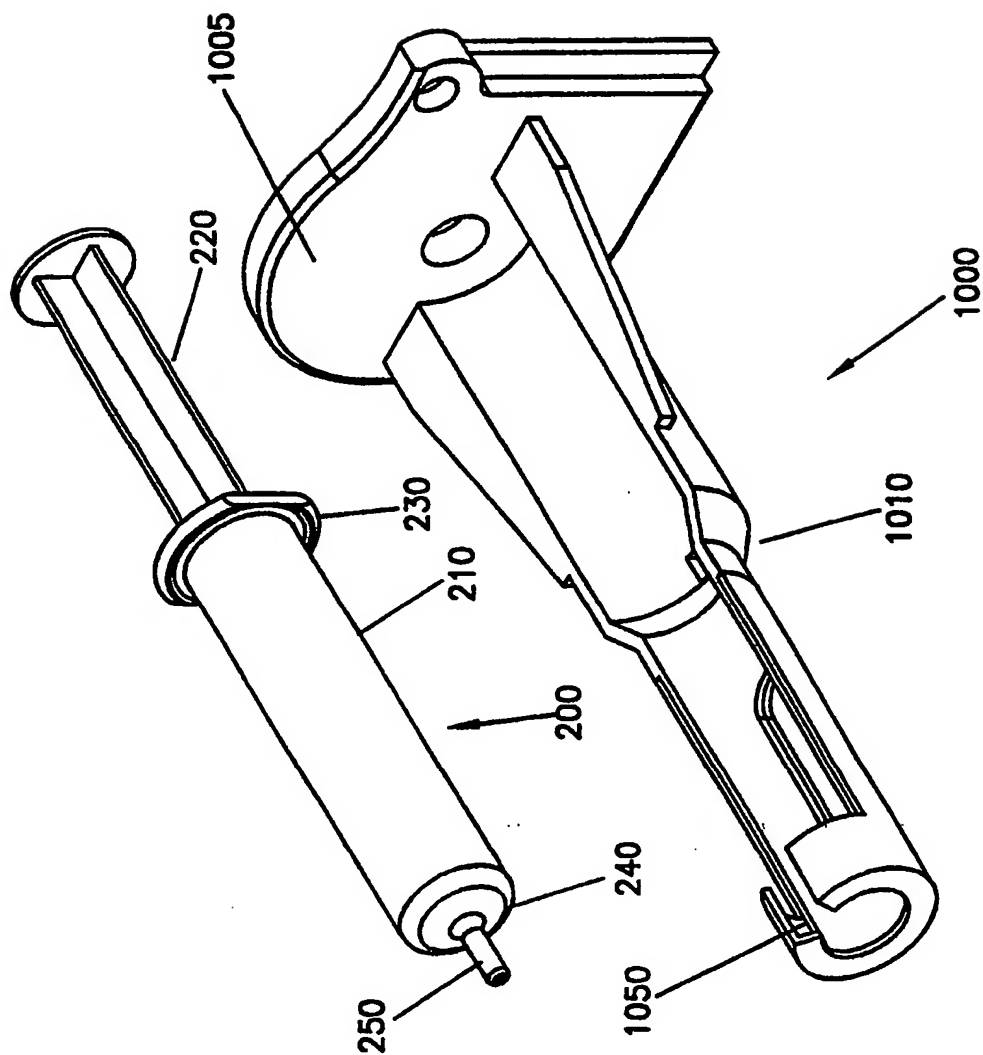


FIG. 8D

33/43

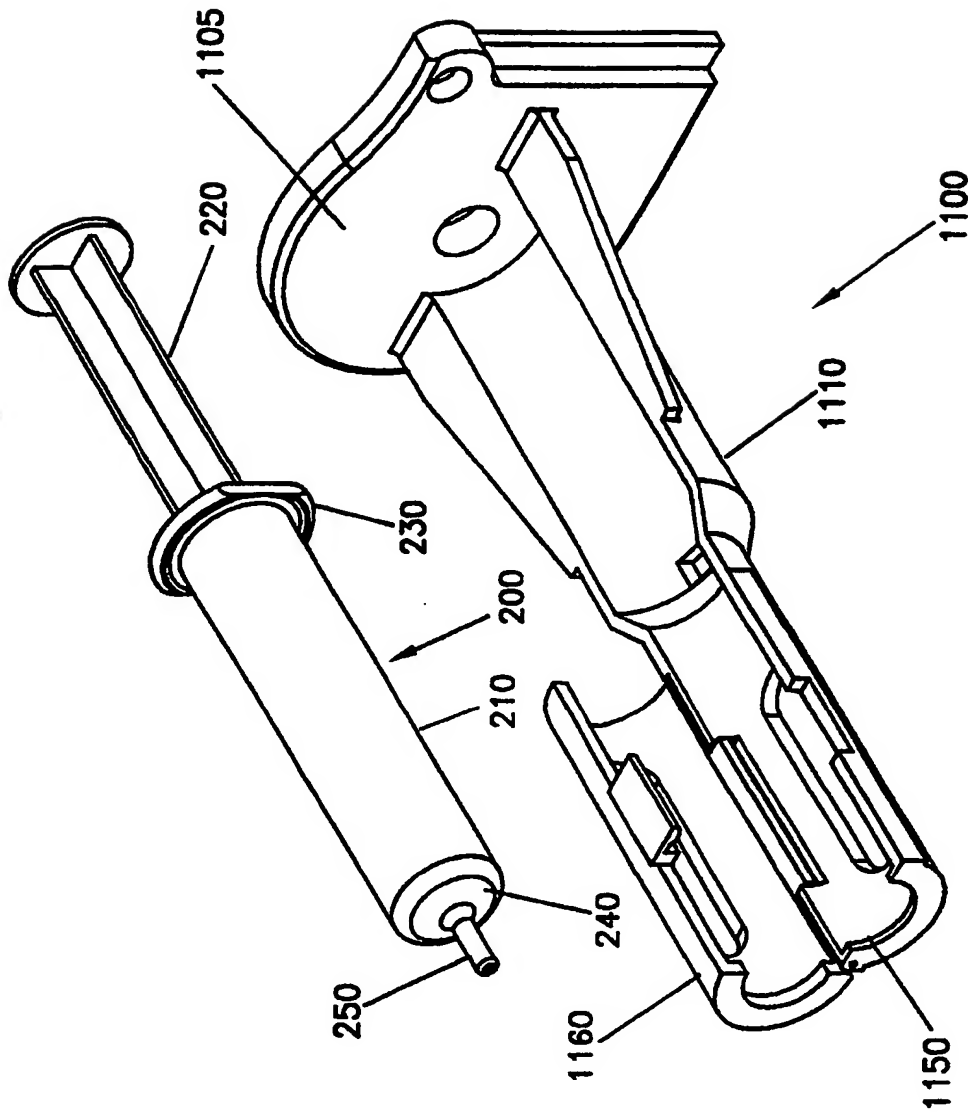


FIG. 8E

34/43

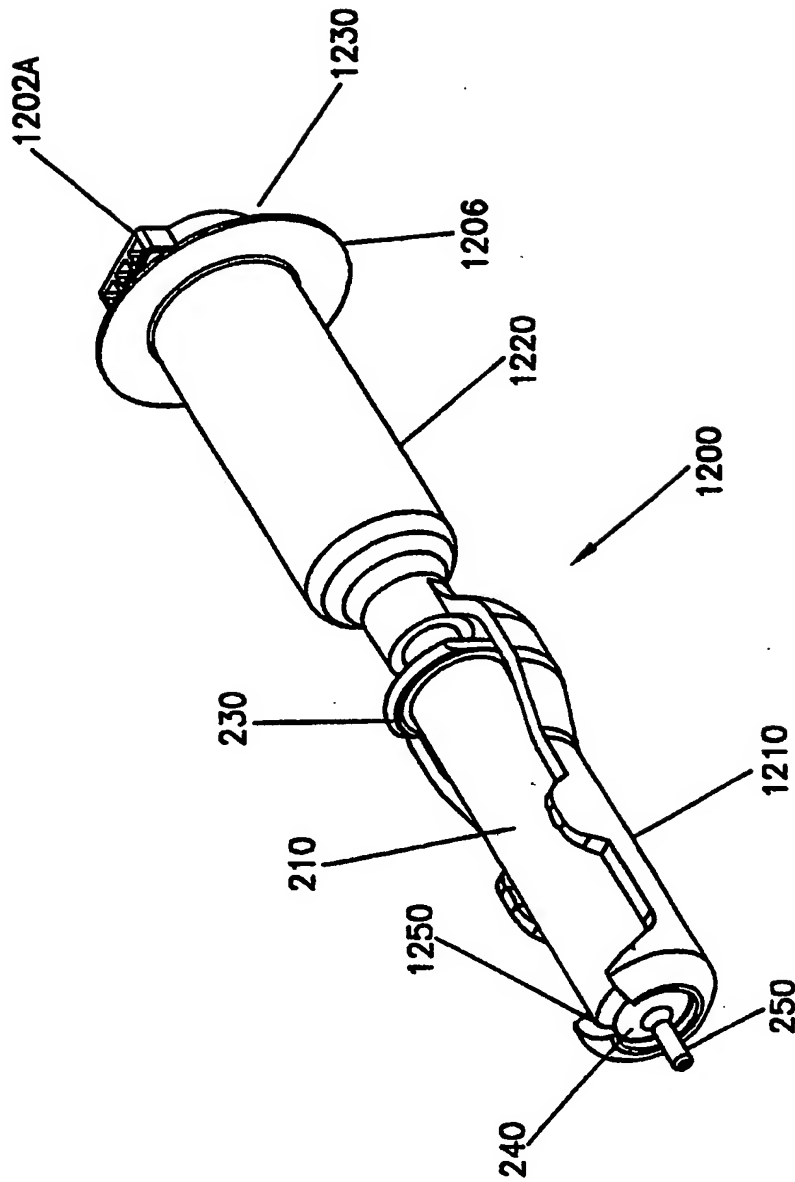


FIG. 9A

35/43

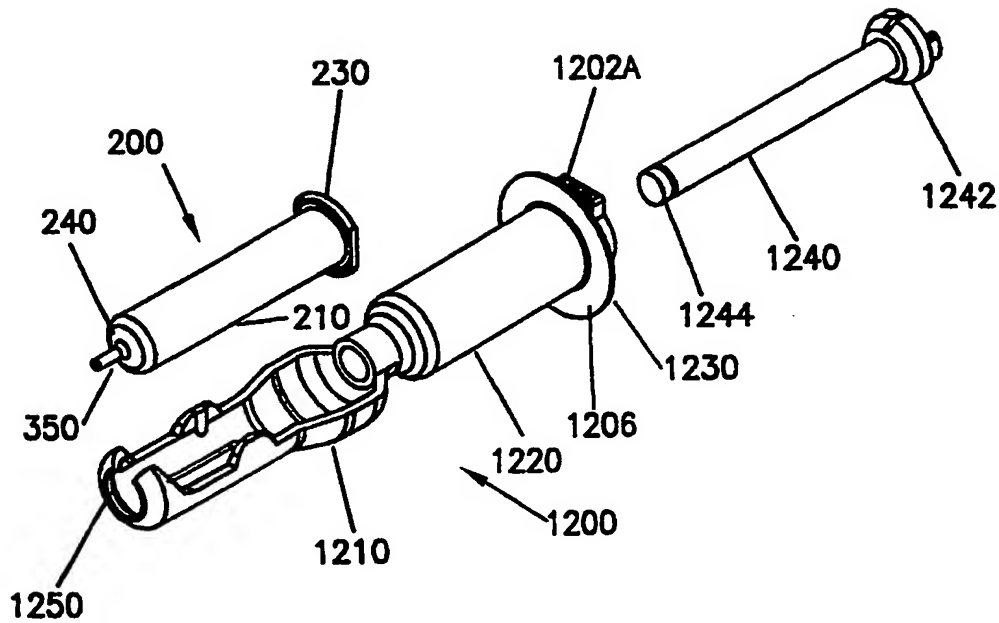


FIG. 9B



36/43

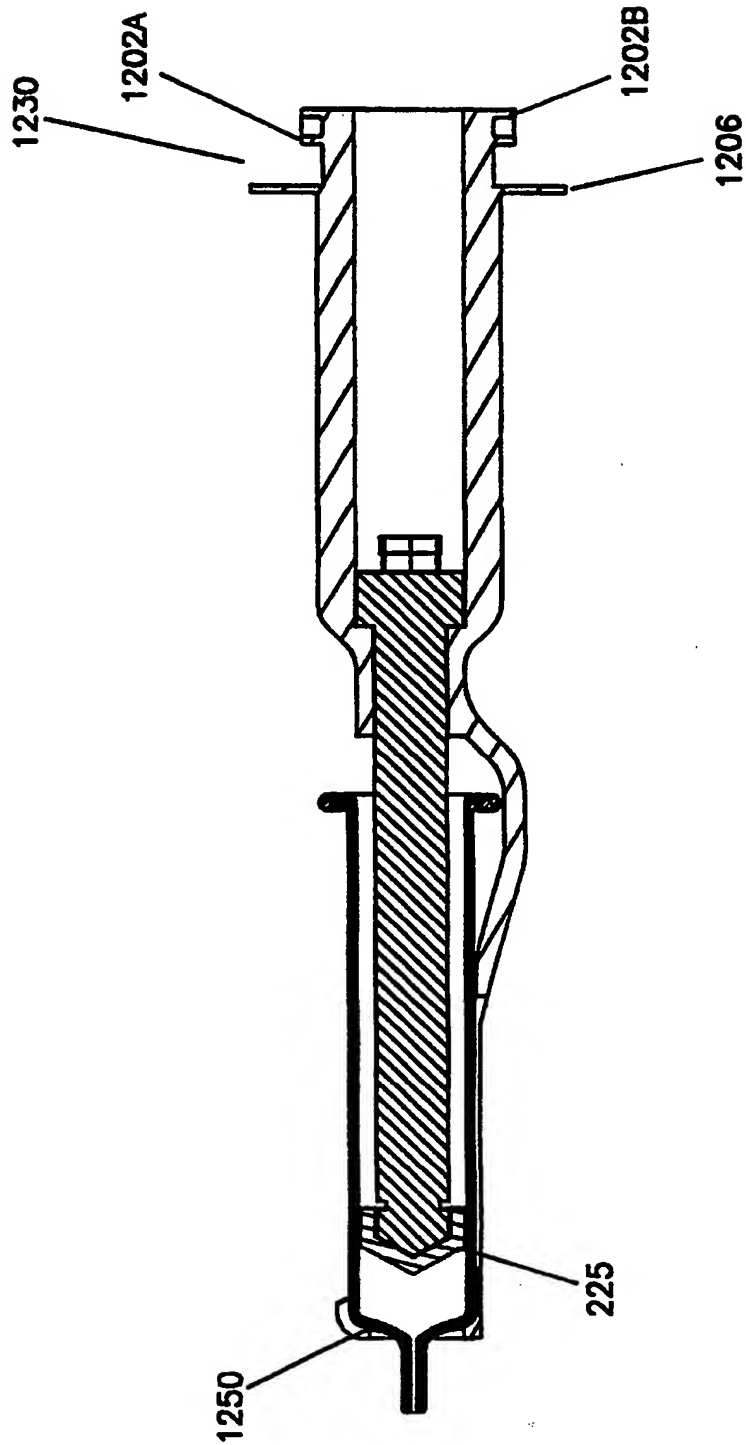


FIG. 9C

37/43

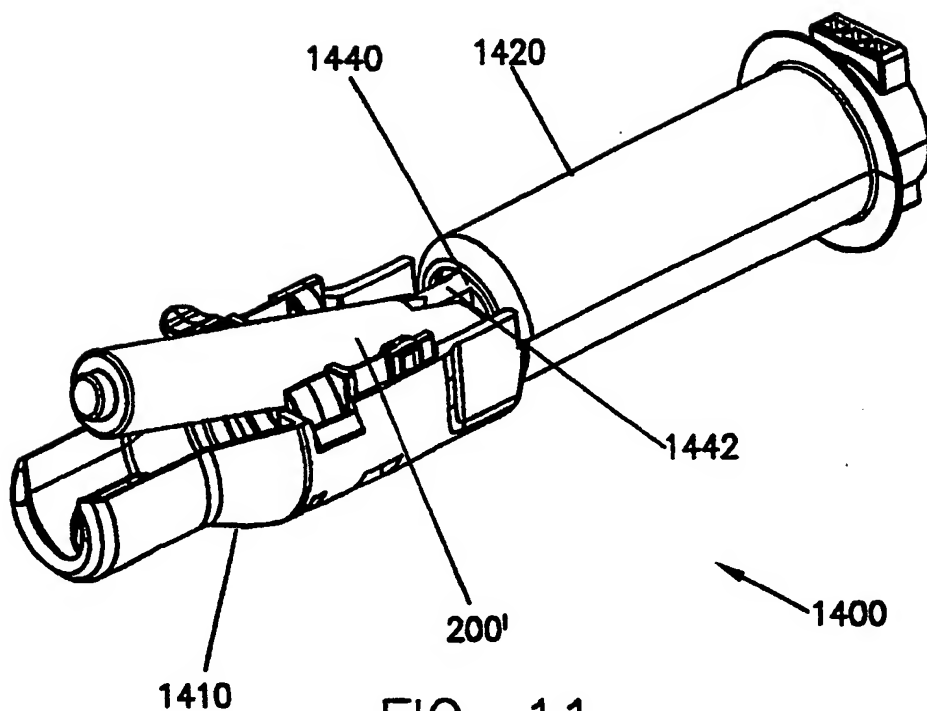


FIG. 11

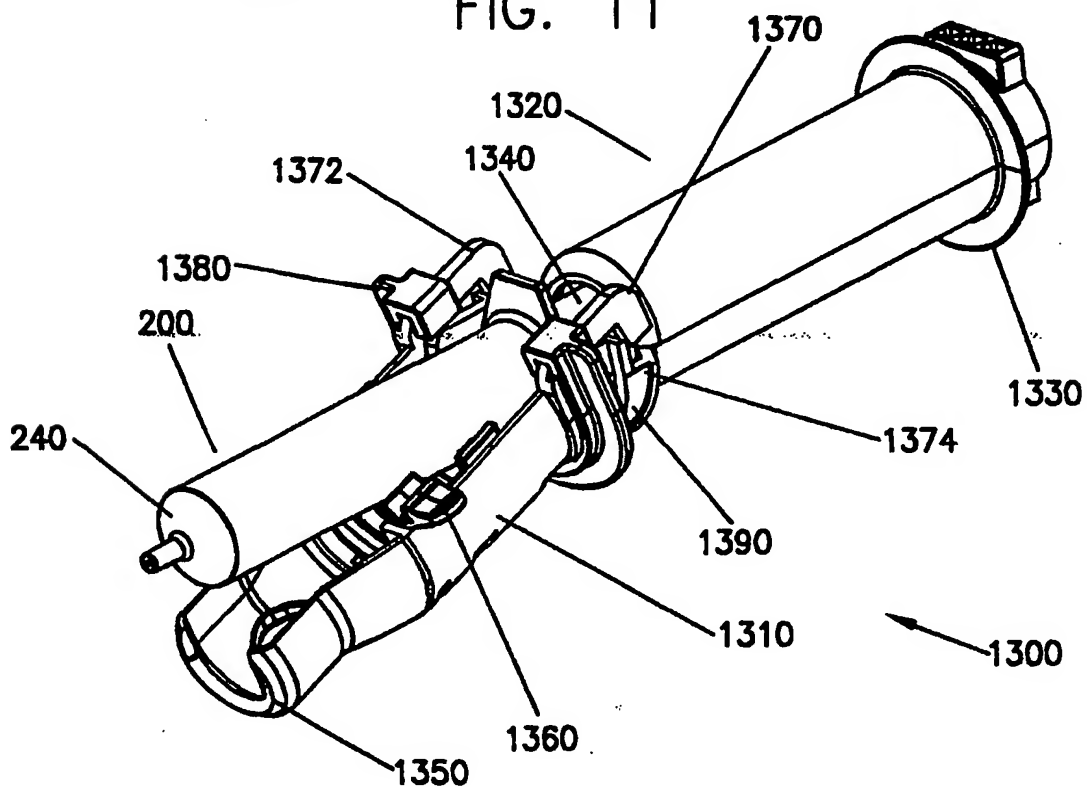


FIG. 10A

38/43

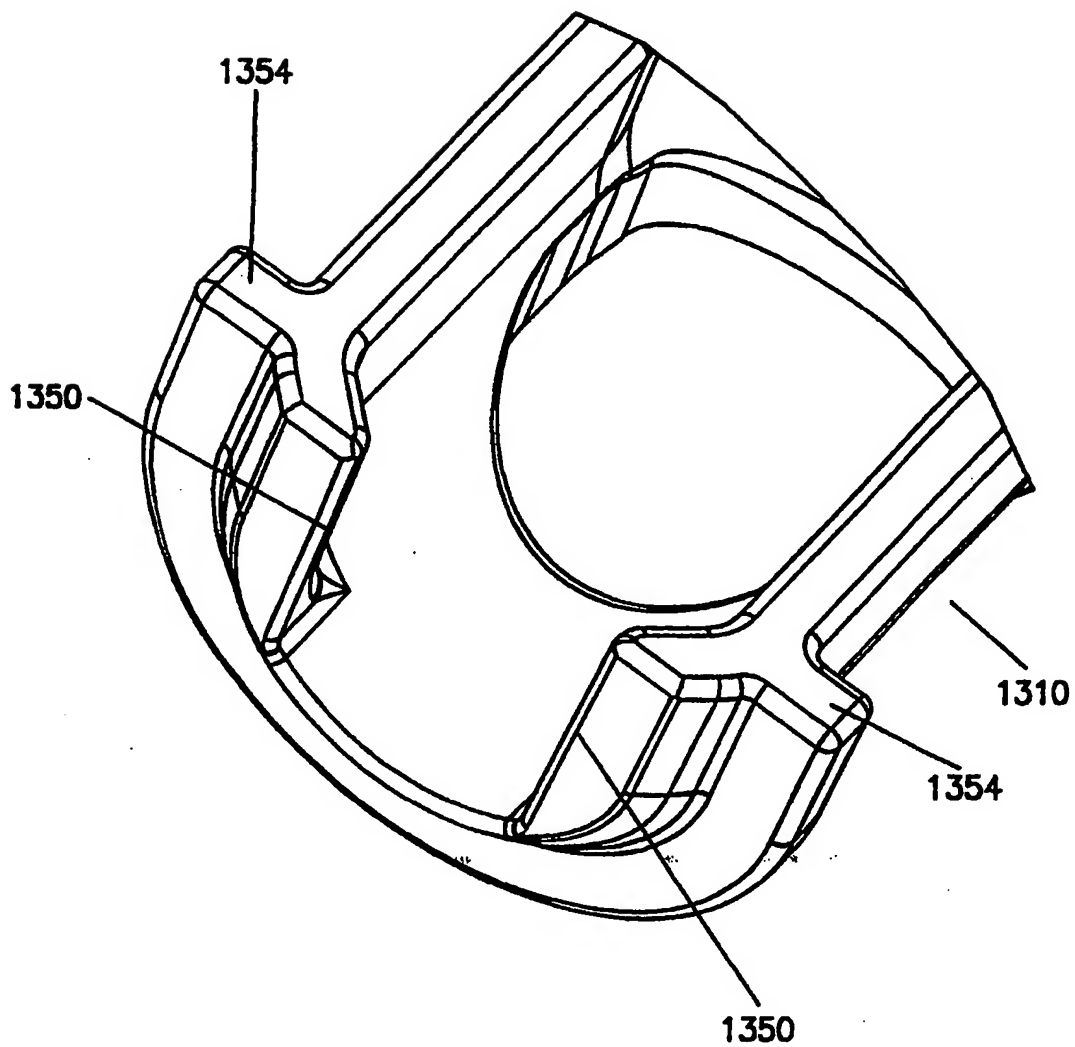


FIG. 10B

39/43

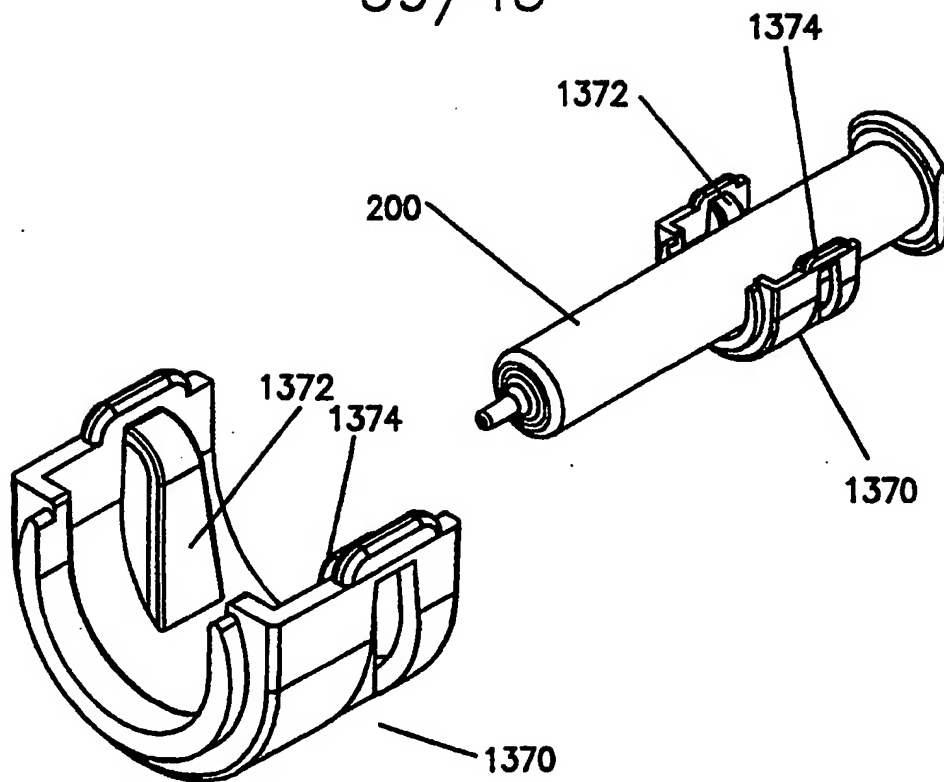


FIG. 10C

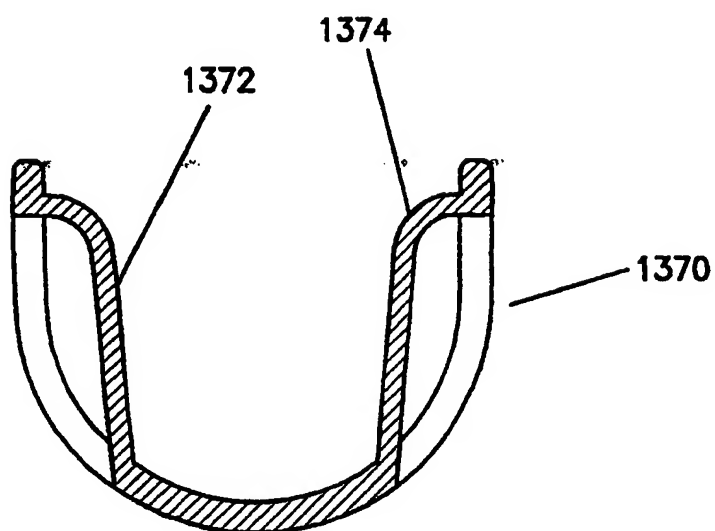


FIG. 10D

40/43

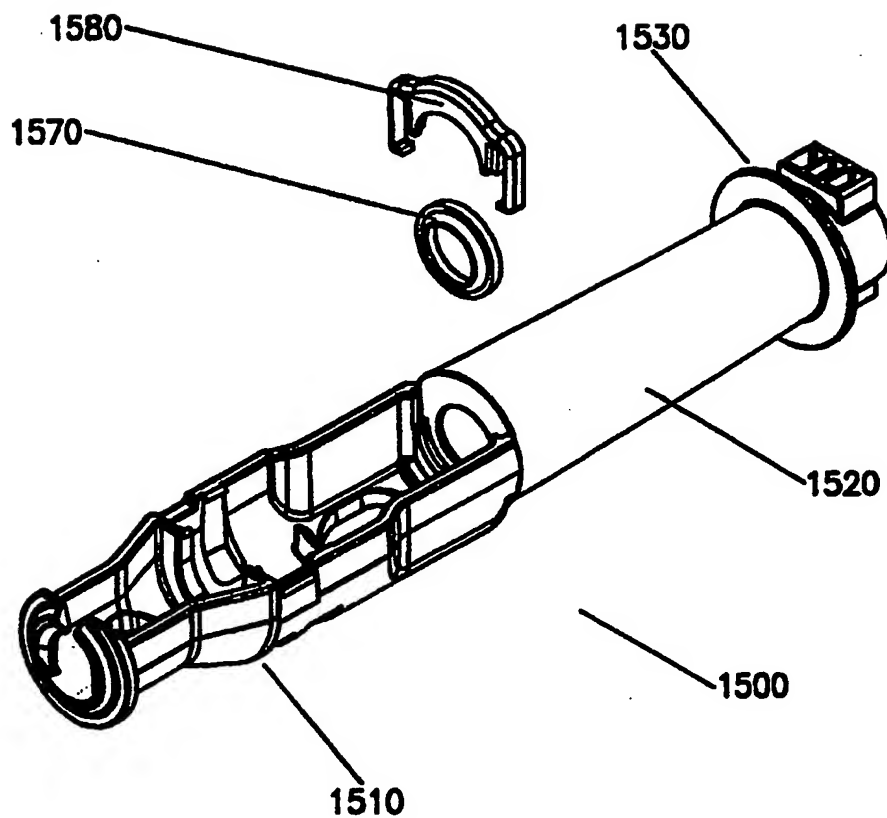


FIG. 12A

41/43

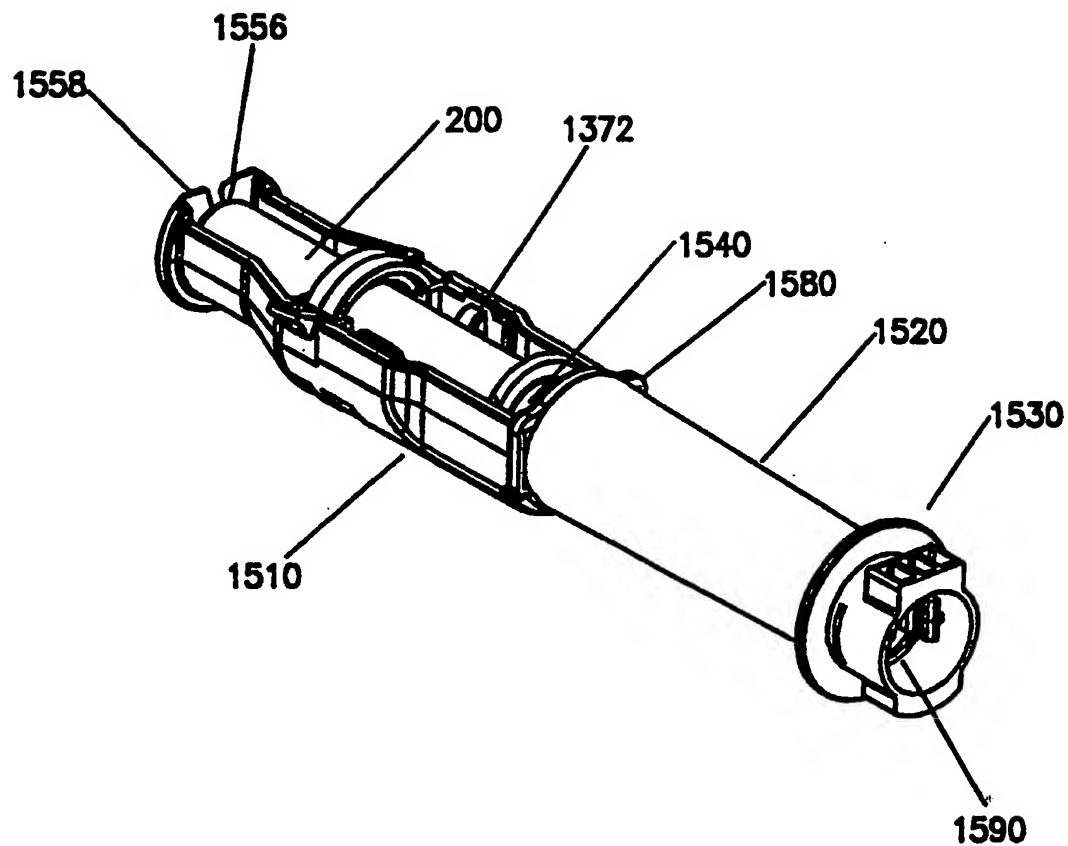


FIG. 12B

42/43

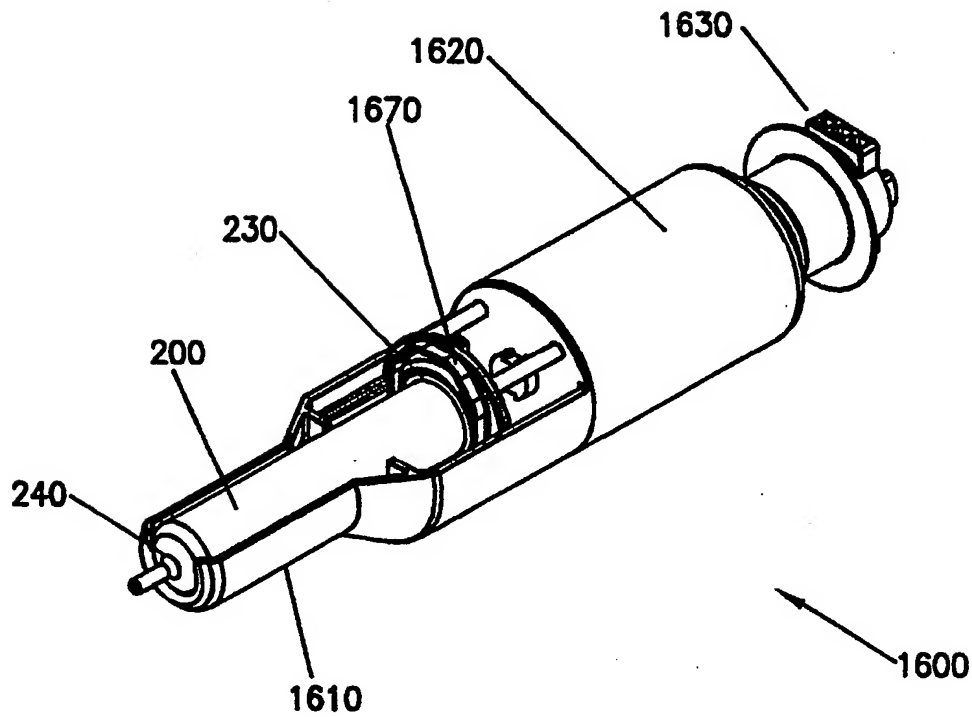


FIG. 13A

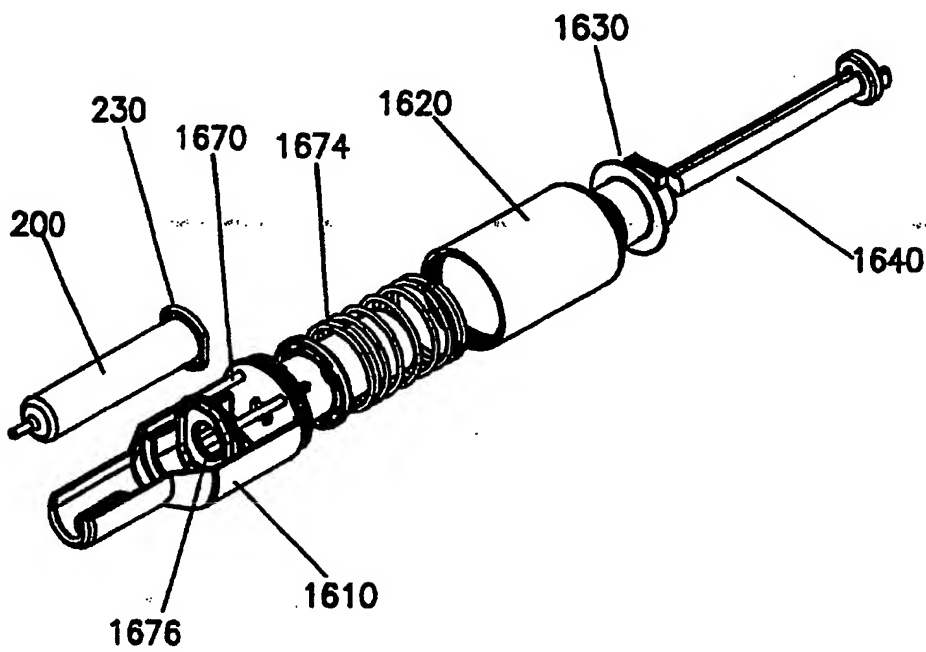


FIG. 13B

43/43

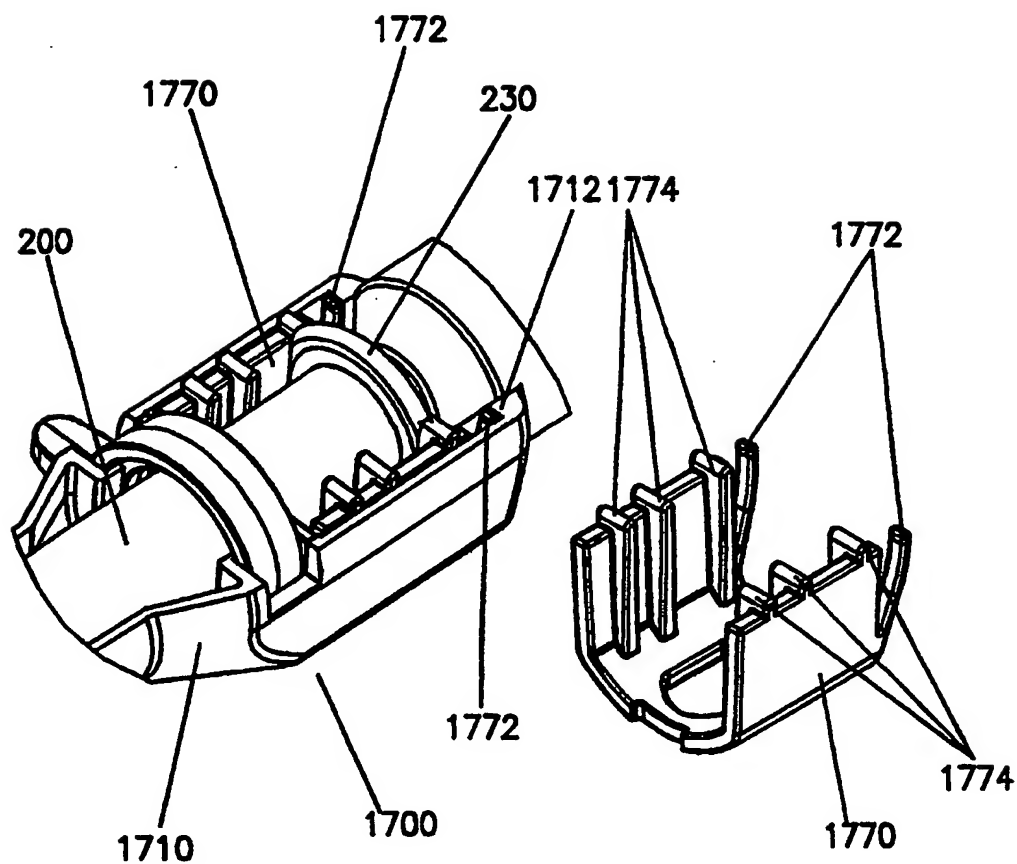


FIG. 14



# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/20623

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 7 A61M5/145

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4 563 175 A (LAFOND MARGARET) 7 January 1986 (1986-01-07) column 8, line 46 - line 55; figures 1,4	8,10
X	US 5 865 805 A (ZIEMBA ROBERT J) 2 February 1999 (1999-02-02) the whole document	1-59
X	US 5 913 844 A (FAGO FRANK M ET AL) 22 June 1999 (1999-06-22) the whole document	1,3,6,8, 9,12-59
X	US 5 520 653 A (REILLY DAVID M ET AL) 28 May 1996 (1996-05-28) cited in the application the whole document	1,3, 13-59
	-/-	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

\* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"8" document member of the same patent family

Date of the actual completion of the international search

15 November 2000

Date of mailing of the international search report

21/11/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3018

Authorized officer

Clarkson, P

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/20623

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 779 675 A (UBER III ARTHUR E ET AL) 14 July 1998 (1998-07-14) the whole document ---	1-59
A	US 5 383 858 A (REILLY DAVID M ET AL) 24 January 1995 (1995-01-24) cited in the application the whole document -----	1-59

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/20623

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4563175 A	07-01-1986	NONE	
US 5865805 A	02-02-1999	JP 11155949 A	15-06-1999
US 5913844 A	22-06-1999	EP 1011760 A WO 9857684 A	28-06-2000 23-12-1998
US 5520653 A	28-05-1996	EP 0847287 A EP 0987040 A JP 11512016 T WO 9709077 A	17-06-1998 22-03-2000 19-10-1999 13-03-1997
US 5779675 A	14-07-1998	AU 6848996 A BR 9610384 A CN 1193918 A EP 0851775 A JP 11511356 T WO 9707841 A US 5899885 A US 5938639 A	19-03-1997 21-12-1999 23-09-1998 08-07-1998 05-10-1999 06-03-1997 04-05-1999 17-08-1999
US 5383858 A	24-01-1995	AT 181676 T CA 2097848 A DE 69325482 D DE 69325482 T DK 584531 T EP 0584531 A EP 0900573 A ES 2134230 T JP 2544571 B JP 6154322 A US 6090064 A US 5795333 A US 5741232 A US 5997502 A	15-07-1999 18-02-1994 05-08-1999 28-10-1999 22-11-1999 02-03-1994 10-03-1999 01-10-1999 16-10-1996 03-06-1994 18-07-2000 18-08-1998 21-04-1998 07-12-1999